

CORRECTED VERSION

(19) World Intellectual Property Organization  
International Bureau(43) International Publication Date  
7 December 2000 (07.12.2000)

PCT

(10) International Publication Number  
WO 00/72827 A3(51) International Patent Classification<sup>7</sup>: A61K 9/16

(21) International Application Number: PCT/US00/14578

(22) International Filing Date: 25 May 2000 (25.05.2000)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:

60/136,323	27 May 1999 (27.05.1999)	US
60/158,659	8 October 1999 (08.10.1999)	US
09/433,486	4 November 1999 (04.11.1999)	US
60/186,310	2 March 2000 (02.03.2000)	US

(81) Designated States (*national*): AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

(71) Applicant: ACUSPHERE, INC. [US/US]; 38 Sidney Street, Cambridge, MA 02139 (US).

(88) Date of publication of the international search report:  
25 January 2001

(72) Inventors: STRAUB, Julie; 100 Cambridge Street, Winchester, MA 01890 (US). BERNSTEIN, Howard; 33A Trowbridge Street, Cambridge, MA 02138 (US). CHICKERING, Donald, E. III; 3 Holly Way, Framingham, MA 01701 (US). KHATAK, Sarwat; 98 Pine Street, Cambridge, MA 02139 (US). RANDALL, Greg; 100 Ledge-wood Drive, Stonham, MA 02180 (US).

(48) Date of publication of this corrected version:  
31 January 2002(15) Information about Correction:  
see PCT Gazette No. 05/2002 of 31 January 2002, Section II

(74) Agents: PABST, Patrea, L. et al.; Holland &amp; Knight, One Atlantic Center, Suite 2000, 1201 West Peachtree Street, Atlanta, GA 30309-3400 (US).

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: POROUS DRUG MATRICES AND METHODS OF MANUFACTURE THEREOF

(57) Abstract: Drugs, especially low aqueous solubility drugs, are provided in a porous matrix form, preferably microparticles, which enhances dissolution of the drug in aqueous media. The drug matrices preferably are made using a process that includes (i) dissolving a drug, preferably a drug having low aqueous solubility, in a volatile solvent to form a drug solution, (ii) combining at least one pore forming agent with the drug solution to form an emulsion, suspension, or second solutions, and (iii) removing the volatile solvent and pore forming agent from the emulsion, suspension, or second solution to yield the porous matrix of drug. The pore forming agent can be either a volatile liquid that is immiscible with the drug solvent or a volatile solid compound, preferably a volatile salt. In a preferred embodiment, spray drying is used to remove the solvents and the pore forming agent. The resulting porous matrix has a faster rate of dissolution following administration to a patient, as compared to non-porous matrix forms of the drug. In a preferred embodiment, microparticles of the porous drug matrix are reconstituted with an aqueous medium and administered parenterally, or processed using standard techniques into tablets or capsules for oral administration. Paclitaxel or docetaxel can be provided in a porous matrix form, which allows the drug to be formulated without solubilizing agents and administered as a bolus, processed using standard techniques into tablets or capsules for oral administration. Paclitaxel or docetaxel can be provided in a porous matrix form, which allows the drug to be formulated without solubilizing agents and administered as a bolus.

WO 00/72827 A3

## POROUS DRUG MATRICES AND METHODS OF MANUFACTURE THEREOF

### Background of the Invention

This invention generally relates to formulations of drugs, especially  
5 drugs having low solubility, and more particularly to methods of making  
formulations of such drugs to enhance their rate of dissolution.

The bioavailability of a drug can be limited by poor dissolution of the  
drug into aqueous bodily fluids following administration. This rate-limiting  
step may therefore be critical to rapidly attaining therapeutically effective  
10 drug levels.

Traditional approaches to parenteral delivery of poorly soluble drugs  
include using large volumes of aqueous diluents, solubilizing agents,  
detergents, non-aqueous solvents, or non-physiological pH solutions. These  
formulations, however, can increase the systemic toxicity of the drug  
15 composition or damage body tissues at the site of administration.

For example, paclitaxel is a natural product which has been shown to  
possess cytotoxic and antitumor activity. While having an unambiguous  
reputation of tremendous therapeutic potential, paclitaxel as a therapeutic  
agent has some patient related drawbacks. These stem, in part, from its  
20 extremely low solubility in water, which makes it difficult to provide in  
suitable dosage form. Because of paclitaxel's poor aqueous solubility, the  
current approved (U.S. FDA) clinical formulation consists of a 6 mg/ml  
solution of paclitaxel in 50% polyoxyethylated castor oil (CREMOPHOR  
EL<sup>TM</sup>) and 50% dehydrated alcohol. *Am. J. Hosp. Pharm.*, 48:1520-24  
25 (1991). In some instances, severe reactions, including hypersensitivity,  
occur in conjunction with the CREMOPHOR<sup>TM</sup> administered in conjunction  
with paclitaxel to compensate for its low water solubility. As a result of the  
incidence of hypersensitivity reactions to the commercial paclitaxel  
formulations and the potential for paclitaxel precipitation in the blood, the  
30 formulation must be infused over several hours. In addition, patients must be  
pretreated with steroids and antihistamines prior to the infusion. In response  
to the hypersensitivity related to the CREMOPHOR<sup>TM</sup>, the increasing  
recognition of paclitaxel's promise as an antineoplastic, and the

undesirability of having to infuse the paclitaxel over several hours, there remains a need to develop improved formulations of the paclitaxel which can be administered as bolus injections. Similarly, it would be advantageous to administer docetaxel by bolus injection, rather than by infusion following  
5 dissolution of the drug in 100% Polysorbate 80, as is the current practice.

Other approaches to parenteral delivery of poorly soluble drugs have focused on the physical form of the drug itself. Since the dissolution rate of a drug particle is directly related to its surface area available to contact the aqueous media at the site of administration or site of absorption, methods of  
10 preparing drugs in nanoparticulate form have been developed in an effort to maximize the drug surface area, as described, for example, in U.S. Patent No. 5,534,270 to De Castro and U.S. Patent No. 5,587,143 to Wong. Nanoparticles, however, can be difficult to produce and maintain in a stable form due to the tendency of the nanoparticles to flocculate or agglomerate,  
15 particularly without the presence of surface modifying agents adsorbed or coated onto the particles. Furthermore, milling or wet grinding techniques, which are typically employed for nanonization, can be undesirable, as it can take several days to process a single batch, scaling-up of the milling or grinding process can be difficult and/or costly, the process can be difficult to  
20 conduct aseptically, and it is difficult to eliminate shedding of milling media into the product.

Other efforts directed at enhancing the rate of dissolution have focused on delivering the drug as a dispersion in a water-soluble or biodegradable matrix, typically in the form of polymeric microparticles. For  
25 example, the dissolution rate of dexamethasone reportedly was improved by entrapping the drug in chitosan microspheres made by spray-drying (Genta, et al., *S.T.P. Pharma Sciences* 5(3):202-07 (1995)). Similarly, others have reported enhanced dissolution rates by mixing a poorly soluble drug powder with a water-soluble gelatin, which purportedly makes the surface of the  
30 drug hydrophilic (Imai, et al., *J. Pharm. Pharmacol.*, 42:615-19 (1990)).

Related efforts have been directed to forming relatively large, porous matrices of low solubility drugs. For example, Roland & Paeratakul, "Spherical Agglomerates of Water-Insoluble Drugs," *J. Pharma. Sci.*,

78(11):964-67 (1989) discloses preparing beads having a low solubility drug content up to 98%, wherein the beads have a porous internal structure. Such large beads, however, are unsuitable for parenteral administration, and the beads have less surface area and slower dissolution rates than smaller particles.

It is therefore an object of the present invention to provide compositions enhancing the dissolution rate of drugs, especially drugs having low aqueous solubility, and to provide methods of making such compositions.

It is another object of the present invention to provide compositions providing enhanced dissolution of drugs, especially drugs of low aqueous solubility, in a formulation suitable for administration by a variety of routes, including, but not limited to, parenteral, mucosal, oral, and topical administration, for local, regional, or systemic effect.

It is another object of the present invention to provide compositions providing enhanced dissolution of paclitaxel in a formulation suitable for administration by a variety of routes, including, but not limited to, parenteral, mucosal, oral, and topical administration, for local, regional, or systemic effect.

It is still another object of the present invention to provide compositions of paclitaxel or docetaxel without the required solubilizing agents present in current commercial formulations.

It is a further object of the present invention to provide compositions, particularly paclitaxel or docetaxel, for administration as a bolus injection instead of by infusion.

### Summary of the Invention

Drugs are provided in a porous matrix form wherein the dissolution rate of the drug is enhanced when the matrix is contacted with an aqueous medium. In a preferred embodiment, low aqueous solubility drugs are provided in a porous matrix form which forms microparticles when the matrix is contacted with an aqueous medium. The porous matrix with low aqueous solubility drugs yields upon contact with an aqueous medium microparticles having a mean diameter between about 0.01 and 5  $\mu\text{m}$  and a

total surface area greater than about  $0.5 \text{ m}^2/\text{mL}$ . The dry porous matrix preferably is in a dry powder form having a TAP density less than or equal to  $1.0 \text{ g/mL}$ .

In a preferred embodiment, the drug is a taxane. For example,  
5 paclitaxel or docetaxel can be provided in a porous matrix form which forms nanoparticles and microparticles of the drug when the matrix is contacted with an aqueous medium.

The porous matrices that contain the drug preferably are made using a process that includes (i) dissolving a drug in a volatile solvent to form a drug  
10 solution, (ii) combining at least one pore forming agent with the drug solution to form an emulsion, suspension, or second solution, and (iii) removing the volatile solvent and pore forming agent from the emulsion, suspension, or second solution to yield the dry porous matrix of drug. The resulting porous matrix has a faster rate of dissolution following  
15 administration to a patient, as compared to non-porous matrix forms of the drug. The pore forming agent can be either a volatile liquid that is immiscible with the drug solvent or a volatile solid compound, preferably a volatile salt. If the pore forming agent is a liquid, the agent is emulsified with the drug solution. If the pore forming agent is a solid, the agent is (i)  
20 dissolved in the drug solution, (ii) dissolved in a solvent that is not miscible in the drug solvent and then emulsified with the drug solution, or (iii) suspended as solid particulates in the drug solution. Optionally, hydrophilic excipients, wetting agents, and/or tonicity agents may be added to the drug solvent, the pore forming agent solvent, or both. The solution, emulsion, or  
25 suspension of the pore forming agent in the drug solution is then processed to remove the drug solvent and the pore forming agent, as well as any pore forming agent solvent. In a preferred embodiment, spray drying, optionally followed by lyophilization, fluid bed drying, or vacuum drying, is used to remove the solvents and the pore forming agent.

30 In a preferred embodiment, the porous drug matrix is reconstituted with an aqueous medium and administered parenterally, such as intramuscularly, subcutaneously, or intravenously. Alternatively, the porous drug matrix can be further processed using standard techniques into tablets or

capsules for oral administration or into rectal suppositories, delivered using a dry powder inhaler for pulmonary administration, or mixed/processed into a cream or ointment for topical administration.

5 An advantage of the porous drug matrix formulations is that they can be administered as a bolus, when the drug, such as paclitaxel, normally must be infused to avoid precipitation of the drug. By avoiding precipitation of drug *in vivo*, the formulations can also be administered intrarterially, intravenously, locally, intracranially, intrathecally, or, if appropriate, directly into a tumor. An additional advantage is the formulations can be  
10 administered in reduced volumes.

In one embodiment, the matrix further includes a pegylated excipient, such as pegylated phospholipid, with the drug. The pegylated excipient shields the drug from macrophage uptake, which prolong its half-life or enhance bioavailability of the drug.

#### 15 **Brief Description of the Drawings**

Figure 1 is a graph of the *in vitro* dissolution rate (percent dissolved versus time) for non-formulated prednisone and prednisone in porous matrix form.

20 Figure 2 is a graph of the *in vitro* dissolution rate (percent dissolved versus time) for non-formulated griseofulvin and griseofulvin in porous matrix form .

Figure 3 is a graph of the *in vitro* dissolution rate (percent dissolved versus time) for non-formulated nifedipine and nifedipine in porous matrix form.

25 Figure 4 is a graph of the *in vitro* dissolution rate (percent dissolved versus time) for non-formulated naproxen and naproxen in a porous matrix form.

Figure 5 is a graph of the *in vitro* dissolution rate (percent dissolved versus time) for non-formulated paclitaxel and paclitaxel in a porous matrix  
30 form.

Figure 6 is a graph of the *in vitro* dissolution rate (percent dissolved versus time) for various porous matrix forms of nifedipine.

Figure 7 is a graph of the *in vitro* dissolution rate (percent dissolved

versus time) for various porous matrix forms of griseofulvin.

Figure 8 is a graph of nifedipine plasma levels versus time post intravenous administration of reconstituted nifedipine matrix in dogs.

Figure 9 shows the chemical structure of taxane compounds,  
5 including paclitaxel and docetaxel.

### Detailed Description of the Invention

The rate of dissolution of drugs can be enhanced by making the drug into a porous matrix form, substantially increasing the surface area of the drug available to contact aqueous biological fluids at the site of  
10 administration of the drug composition. The method for making the porous matrix of drug includes the steps of (a) dissolving a drug in a volatile solvent to form a drug solution, (b) combining at least one pore forming agent with the drug solution to form an emulsion, suspension, or second solution, and (c) removing the volatile solvent and pore forming agent from the emulsion,  
15 suspension, or second solution to yield the porous matrix of drug.

In a preferred embodiment, pharmaceutical compositions are provided for paclitaxel, or docetaxel, without the solubilizing agent CREMOPHOR<sup>TM</sup>, or Polysorbate 80, respectively, wherein the pharmaceutical composition can be administered as a bolus. The  
20 compositions are porous dry powders, which upon the addition of an aqueous medium form a suspension of nanoparticles and microparticles of the drug.

#### I. Drug Matrix Compositions

The porous drug matrix is at least 1 to 95%, preferably at least about 10%, and more preferably between about 10 and 70%, drug by weight. The  
25 matrices also may contain hydrophilic excipients such as water soluble polymers or sugars, wetting agents such as surfactants, and tonicity agents.

The form of the drug matrix (drug powder) is critical to the dissolution rate. The matrix must contain microparticles of drug, which preferably have a mean diameter between about 0.01 and 5  $\mu\text{m}$ , more  
30 preferably between about 0.1 and 5  $\mu\text{m}$ . In one embodiment, the mean diameter of the microparticles is between about 1 and 5  $\mu\text{m}$ . The average total surface area of the microparticles contained within the porous matrix, which typically is in the form of a dry powder, is 0.5  $\text{m}^2/\text{mL}$  or greater,

preferably  $0.9 \text{ m}^2/\text{mL}$  or greater. Total surface area values for the microparticles can be determined using standard Coulter Counter equipment and techniques.

5 The drug matrix must be sufficiently porous to yield, upon contact with an aqueous medium, microparticles having these parameters. Measurements useful in characterizing the porosity of the drug matrix are the bulk density or the transaxial pressure ("TAP") density of the dry porous matrix (dry powder) and the total surface area (sum of internal and external surface area) of the dry porous matrix. The TAP density preferably is less than or equal to  $1.0 \text{ g/ml}$ , more preferably less than  $0.8 \text{ g/ml}$ . This level of porosity of the matrix, characterized by density, provides sufficient surface area to enhance wetting of the dry porous matrix and enhance drug dissolution. The total surface area of the porous matrix can be measured, for example, by BET surface area analysis. In some embodiments, the total surface area of the porous matrix preferably is greater than  $0.1 \text{ m}^2/\text{g}$ , more preferably greater than or equal to  $0.2 \text{ m}^2/\text{g}$ . This level of total surface area provides sufficient surface area to enhance wetting of the dry porous matrix and enhance drug dissolution.

### 1. Drugs

20 A wide variety of drugs are useful in the methods and compositions described herein. In a preferred embodiment, the drug is a low aqueous solubility drug. As used herein, the term "low aqueous solubility" means that the drug has a solubility of less than about  $10 \text{ mg/mL}$ , and preferably less than about  $5 \text{ mg/mL}$ , in aqueous media at approximately physiological temperatures and pH. As used herein, the term "drug" refers to chemical or biological molecules providing a therapeutic, diagnostic, or prophylactic effect *in vivo*. A particularly preferred class of drugs is taxanes.

#### (i) Taxanes

25 Taxanes are anticancer cytotoxics that stabilize cellular microtubules. Taxane compounds useful in the compositions and methods described herein include paclitaxel and docetaxel, as well as natural and synthetic analogs thereof which possess anticancer or anti-angiogenic activity. Paclitaxel and docetaxel have substantial activity, and one or both of these agents are



widely accepted as components of therapy for advanced breast, lung, and ovarian carcinomas. Paclitaxel has recently been approved in the USA for the adjuvant treatment of early stage node-positive breast carcinoma. The chemical structure of taxanes, including paclitaxel and docetaxel, is shown in

5 Figure 9.

(ii) Other Drugs

Other drugs contemplated for use in the compositions described herein include the following categories and examples of drugs and alternative forms of these drugs such as alternative salt forms, free acid

10 forms, free base forms, and hydrates:

analgesics/antipvretics (e.g., aspirin, acetaminophen, ibuprofen, naproxen sodium, buprenorphine, propoxyphene hydrochloride, propoxyphene napsylate, meperidine hydrochloride, hydromorphone hydrochloride, morphine, oxycodone, codeine, dihydrocodeine bitartrate, pentazocine,

15 hydrocodone bitartrate, levorphanol, diflunisal, trolamine salicylate, nalbuphine hydrochloride, mefenamic acid, butorphanol, choline salicylate, butalbital, phenyltoloxamine citrate, diphenhydramine citrate, methotrimeprazine, cinnamedrine hydrochloride, and meprobamate);

antiasthmatics (e.g., ketotifen and traxanox);

20 antibiotics (e.g., neomycin, streptomycin, chloramphenicol, cephalosporin, ampicillin, penicillin, tetracycline, and ciprofloxacin);

antidepressants (e.g., nefopam, oxypertine, doxepin, amoxapine, trazodone, amitriptyline, maprotiline, phenelzine, desipramine, nortriptyline, tranlycypromine, fluoxetine, doxepin, imipramine, imipramine pamoate,

25 isocarboxazid, trimipramine, and protriptyline);

antidiabetics (e.g., biguanides and sulfonylurea derivatives);

antifungal agents (e.g., griseofulvin, ketoconazole, itraconazole, amphotericin B, nystatin, and candicidin);

antihypertensive agents (e.g., propranolol, propafenone, oxyprenolol,

30 nifedipine, reserpine, trimethaphan, phenoxybenzamine, pargyline hydrochloride, deserpidine, diazoxide, guanethidine monosulfate, minoxidil, rescinnamine, sodium nitroprusside, rauwolfia serpentina, alseroxylon, and phentolamine );

- anti-inflammatories (e.g., (non-steroidal) indomethacin, ketoprofen, flurbiprofen, naproxen, ibuprofen, ramifenazone, piroxicam, (steroidal) cortisone, dexamethasone, fluazacort, celecoxib, rofecoxib, hydrocortisone, prednisolone, and prednisone);
- 5 antineoplastics (e.g., cyclophosphamide, actinomycin, bleomycin, daunorubicin, doxorubicin, epirubicin, mitomycin, methotrexate, fluorouracil, carboplatin, carmustine (BCNU), methyl-CCNU, cisplatin, etoposide, camptothecin and derivatives thereof, phenesterine, vinblastine, vincristine, tamoxifen, and pipsulfan);
- 10 antianxiety agents (e.g., lorazepam, buspirone, prazepam, chlordiazepoxide, oxazepam, clorazepate dipotassium, diazepam, hydroxyzine pamoate, hydroxyzine hydrochloride, alprazolam, droperidol, halazepam, chlormezanone, and dantrolene);
- immunosuppressive agents (e.g., cyclosporine, azathioprine, mizoribine, and
- 15 FK506 (tacrolimus));
- antimigraine agents (e.g., ergotamine, propanolol, isometheptene mucate, and dichloralphenazone);
- sedatives/hypnotics (e.g., barbiturates such as pentobarbital, pentobarbital, and secobarbital; and benzodiazepines such as flurazepam hydrochloride,
- 20 triazolam, and midazolam);
- antianginal agents (e.g., beta-adrenergic blockers; calcium channel blockers such as nifedipine, and diltiazem; and nitrates such as nitroglycerin, isosorbide dinitrate, pentaerythritol tetranitrate, and erythrityl tetranitrate);
- antipsychotic agents (e.g., haloperidol, loxapine succinate, loxapine
- 25 hydrochloride, thioridazine, thioridazine hydrochloride, thiothixene, fluphenazine, fluphenazine decanoate, fluphenazine enanthate, trifluoperazine, chlorpromazine, perphenazine, lithium citrate, and prochlorperazine);
- antimanic agents (e.g., lithium carbonate);
- 30 antiarrhythmics (e.g., bretylium tosylate, esmolol, verapamil, amiodarone, encainide, digoxin, digitoxin, mexiletine, disopyramide phosphate, procainamide, quinidine sulfate, quinidine gluconate, quinidine polygalacturonate, flecainide acetate, tocainide, and lidocaine);

- antiarthritic agents (e.g., phenylbutazone, sulindac, penicillamine, salsalate, piroxicam, azathioprine, indomethacin, meclofenamate, gold sodium thiomalate, ketoprofen, auranofin, aurothioglucose, and tolmetin sodium);
- antigout agents (e.g., colchicine, and allopurinol);
- 5 anticoagulants (e.g., heparin, heparin sodium, and warfarin sodium);
- thrombolytic agents (e.g., urokinase, streptokinase, and alteplase);
- antifibrinolytic agents (e.g., aminocaproic acid);
- hemorheologic agents (e.g., pentoxifylline);
- antiplatelet agents (e.g., aspirin);
- 10 anticonvulsants (e.g., valproic acid, divalproex sodium, phenytoin, phenytoin sodium, clonazepam, primidone, phenobarbital, carbamazepine, amobarbital sodium, methsuximide, metharbital, mephobarbital, mephentyoin, phensuximide, paramethadione, ethotoin, phenacemide, secobarbital sodium, clorazepate dipotassium, and trimethadione);
- 15 antiparkinson agents (e.g., ethosuximide);
- antihistamines/antipruritics (e.g., hydroxyzine, diphenhydramine, chlorpheniramine, brompheniramine maleate, cyproheptadine hydrochloride, terfenadine, clemastine fumarate, triprolidine, carbinoxamine, diphenylpyraline, phenindamine, azatadine, tripeleminamine,
- 20 dexchlorpheniramine maleate, and methdilazine);
- agents useful for calcium regulation (e.g., calcitonin, and parathyroid hormone);
- antibacterial agents (e.g., amikacin sulfate, aztreonam, chloramphenicol, chloramphenicol palmitate, ciprofloxacin, clindamycin, clindamycin
- 25 palmitate, clindamycin phosphate, metronidazole, metronidazole hydrochloride, gentamicin sulfate, lincomycin hydrochloride, tobramycin sulfate, vancomycin hydrochloride, polymyxin B sulfate, colistimethate sodium, and colistin sulfate);
- antiviral agents (e.g., interferon alpha, beta or gamma, zidovudine,
- 30 amantadine hydrochloride, ribavirin, and acyclovir);
- antimicrobials (e.g., cephalosporins such as cefazolin sodium, cephadrine, cefaclor, cephapirin sodium, ceftizoxime sodium, cefoperazone sodium, cefotetan disodium, cefuroxime e azotil, cefotaxime sodium, cefadroxil

- monohydrate, cephalixin, cephalothin sodium, cephalixin hydrochloride monohydrate, cefamandole nafate, cefoxitin sodium, cefonicid sodium, ceforanide, ceftriaxone sodium, ceftazidime, cefadroxil, cephradine, and cefuroxime sodium; penicillins such as ampicillin, amoxicillin, penicillin G
- 5 benzathine, cyclacillin, ampicillin sodium, penicillin G potassium, penicillin V potassium, piperacillin sodium, oxacillin sodium, bacampicillin hydrochloride, cloxacillin sodium, ticarcillin disodium, azlocillin sodium, carbenicillin indanyl sodium, penicillin G procaine, methicillin sodium, and nafcillin sodium; erythromycins such as erythromycin ethylsuccinate,
- 10 erythromycin, erythromycin estolate, erythromycin lactobionate, erythromycin stearate, and erythromycin ethylsuccinate; and tetracyclines such as tetracycline hydrochloride, doxycycline hyclate, and minocycline hydrochloride, azithromycin, clarithromycin);
- anti-infectives (e.g., GM-CSF);
- 15 bronchodilators (e.g., sympathomimetics such as epinephrine hydrochloride, metaproterenol sulfate, terbutaline sulfate, isoetharine, isoetharine mesylate, isoetharine hydrochloride, albuterol sulfate, albuterol, bitolterolmesylate, isoproterenol hydrochloride, terbutaline sulfate, epinephrine bitartrate, metaproterenol sulfate, epinephrine, and epinephrine bitartrate;
- 20 anticholinergic agents such as ipratropium bromide; xanthines such as aminophylline, dyphylline, metaproterenol sulfate, and aminophylline; mast cell stabilizers such as cromolyn sodium; inhalant corticosteroids such as beclomethasone dipropionate (BDP), and beclomethasone dipropionate monohydrate; salbutamol; ipratropium bromide; budesonide; ketotifen;
- 25 salmeterol; xinafoate; terbutaline sulfate; triamcinolone; theophylline; nedocromil sodium; metaproterenol sulfate; albuterol; flunisolide; fluticasone propionate;
- steroidal compounds and hormones (e.g., androgens such as danazol, testosterone cypionate, fluoxymesterone, ethyltestosterone, testosterone
- 30 enanthate, methyltestosterone, fluoxymesterone, and testosterone cypionate; estrogens such as estradiol, estropipate, and conjugated estrogens; progestins such as methoxyprogesterone acetate, and norethindrone acetate; corticosteroids such as triamcinolone, betamethasone, betamethasone sodium

- phosphate, dexamethasone, dexamethasone sodium phosphate, dexamethasone acetate, prednisone, methylprednisolone acetate suspension, triamcinolone acetonide, methylprednisolone, prednisolone sodium phosphate, methylprednisolone sodium succinate, hydrocortisone sodium succinate, triamcinolone hexacetonide, hydrocortisone, hydrocortisone cypionate, prednisolone, fludrocortisone acetate, paramethasone acetate, prednisolone tebutate, prednisolone acetate, prednisolone sodium phosphate, and hydrocortisone sodium succinate; and thyroid hormones such as levothyroxine sodium);
- 10 hypoglycemic agents (e.g., human insulin, purified beef insulin, purified pork insulin, glyburide, chlorpropamide, glipizide, tolbutamide, and tolazamide);
- hypolipidemic agents (e.g., clofibrate, dextrothyroxine sodium, probucol, pravastatin, atorvastatin, lovastatin, and niacin);
- 15 proteins (e.g., DNase, alginase, superoxide dismutase, and lipase);
- nucleic acids (e.g., sense or anti-sense nucleic acids encoding any therapeutically useful protein, including any of the proteins described herein);
- agents useful for erythropoiesis stimulation (e.g., erythropoietin);
- 20 antiulcer/antireflux agents (e.g., famotidine, cimetidine, and ranitidine hydrochloride);
- antinauseants/antiemetics (e.g., meclizine hydrochloride, nabilone, prochlorperazine, dimenhydrinate, promethazine hydrochloride, thiethylperazine, and scopolamine);
- 25 oil-soluble vitamins (e.g., vitamins A, D, E, K, and the like);
- as well as other drugs such as mitotane, halonitrosoureas, anthrocyclines, and ellipticine. A description of these and other classes of useful drugs and a listing of species within each class can be found in Martindale, *The Extra Pharmacopoeia*, 30th Ed. (The Pharmaceutical Press, London 1993).
- 30 Examples of other drugs useful in the compositions and methods described herein include ceftriaxone, ketoconazole, ceftazidime, oxaprozin, albuterol, valacyclovir, urofollitropin, famciclovir, flutamide, enalapril, mefformin, itraconazole, buspirone, gabapentin, fosinopril, tramadol,

acarbose, lorazepam, follitropin, glipizide, omeprazole, fluoxetine, lisinopril, tramadol, levofloxacin, zafirlukast, interferon, growth hormone, interleukin, erythropoietin, granulocyte stimulating factor, nizatidine, bupropion, perindopril, erbumine, adenosine, alendronate, alprostadil, benazepril,

5 betaxolol, bleomycin sulfate, dexfenfluramine, diltiazem, fentanyl, flecainid, gemcitabine, glatiramer acetate, granisetron, lamivudine, mangafodipir trisodium, mesalamine, metoprolol fumarate, metronidazole, miglitol, moexipril, monteleukast, octreotide acetate, olopatadine, paricalcitol, somatropin, sumatriptan succinate, tacrine, verapamil, nabumetone,

10 trovafloxacin, dolasetron, zidovudine, finasteride, tobramycin, isradipine, tolcapone, enoxaparin, fluconazole, lansoprazole, terbinafine, pamidronate, didanosine, diclofenac, cisapride, venlafaxine, troglitazone, fluvastatin, losartan, imiglucerase, donepezil, olanzapine, valsartan, fexofenadine, calcitonin, and ipratropium bromide. These drugs are generally considered

15 to be water soluble.

Preferred drugs include albuterol or albuterol sulfate, adapalene, doxazosin mesylate, mometasone furoate, ursodiol, amphotericin, enalapril maleate, felodipine, nefazodone hydrochloride, valrubicin, albendazole, conjugated estrogens, medroxyprogesterone acetate, nicardipine

20 hydrochloride, zolpidem tartrate, amlodipine besylate, ethinyl estradiol, omeprazole, rubitecan, amlodipine besylate/ benazepril hydrochloride, etodolac, paroxetine hydrochloride, atovaquone, felodipine, podofilox, paricalcitol, betamethasone dipropionate, fentanyl, pramipexole dihydrochloride, Vitamin D<sub>3</sub> and related analogues, finasteride, quetiapine

25 fumarate, alprostadil, candesartan, cilexetil, fluconazole, ritonavir, busulfan, carbamazepine, flumazenil, risperidone, carbamazepine, carbidopa, levodopa, ganciclovir, saquinavir, amprenavir, carboplatin, glyburide, sertraline hydrochloride, rofecoxib carvedilol, halobetasolpropionate, sildenafil citrate, celecoxib, chlorthalidone, imiquimod, simvastatin,

30 citalopram, ciprofloxacin, irinotecan hydrochloride, sparfloxacin, efavirenz, cisapride monohydrate, lansoprazole, tamsulosin hydrochloride, mofafinil, clarithromycin, letrozole, terbinafine hydrochloride, rosiglitazone maleate, diclofenac sodium, lomefloxacin hydrochloride, tirofiban hydrochloride,

telmisartan, diazepam, loratadine, toremifene citrate, thalidomide, dinoprostone, mefloquine hydrochloride, trandolapril, docetaxel, mitoxantrone hydrochloride, tretinoin, etodolac, triamcinolone acetate, estradiol, ursodiol, nelfinavir mesylate, indinavir, beclomethasone dipropionate, oxaprozin, flutamide, famotidine, nifedipine, prednisone, 5 cefuroxime, lorazepam, digoxin, lovastatin, griseofulvin, naproxen, ibuprofen, isotretinoin, tamoxifen citrate, nimodipine, amiodarone, and alprazolam.

## 2. Excipients

10 The matrices may contain hydrophilic excipients such as water soluble polymers or sugars which can serve as bulking agents or as wetting agents, wetting agents such as surfactants or sugars, and tonicity agents. Upon contact with an aqueous medium, water penetrates through the highly porous matrix to dissolve the water soluble excipients in the matrix. In the 15 case of low aqueous solubility drugs, a suspension of drug particles in the aqueous medium remains. The total surface area of the resultant low aqueous solubility drug microparticles is increased relative to the unprocessed drug and the dissolution rate of the drug is increased.

One of skill in the art can select appropriate excipients for use in the 20 drug matrix compositions, considering a variety of factors, such as the drug to be administered, the route of administration, the dosage, and the preferred dissolution rate. For example, the excipients can function as bulking agents, release-modifiers, wetting agents, tonicity agents, or combinations thereof. Preferred excipients include hydrophilic polymers, wetting agents, and 25 sugars. The amount of excipient in the drug matrix is less than about 95%, more preferably less than about 80%, by weight of the drug matrix.

The hydrophilic excipients, wetting agents, and tonicity agents may be added to the drug solution, the pore forming agent, or both, during production of the matrix.

### 30 (i) Hydrophilic Polymers

The polymers that can be used in the drug matrices described herein include both synthetic and natural polymers, either non-biodegradable or biodegradable. Representative synthetic polymers include polyethylene

glycol ("PEG"), polyvinyl pyrrolidone, polymethacrylates, polylysine, poloxamers, polyvinyl alcohol, polyacrylic acid, polyethylene oxide, and polyethyloxazoline. Representative natural polymers include albumin, alginate, gelatin, acacia, chitosan, cellulose dextran, ficoll, starch, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxy-propylmethyl cellulose, hyaluronic acid, carboxyethyl cellulose, carboxymethyl cellulose, deacetylated chitosan, dextran sulfate, and derivatives thereof. Preferred hydrophilic polymers include PEG, polyvinyl pyrrolidone, poloxamers, hydroxypropyl cellulose, and hydroxyethyl cellulose.

10 The hydrophilic polymer selected for use in a particular drug matrix formulation is based on a variety of factors, such as the polymer molecular weight, polymer hydrophilicity, and polymer inherent viscosity. The hydrophilic polymer can be used as a bulking agent or as a wetting agent.

(ii) Sugars

15 Representative sugars that can be used in the drug matrices include mannitol, sorbitol, xylitol, glucitol, ducitol, inositol, arabinitol, arabitol, galactitol, iditol, allitol, fructose, sorbose, glucose, xylose, trehalose, allose, dextrose, altrose, gulose, idose, galactose, talose, ribose, arabinose, xylose, lyxose, sucrose, maltose, lactose, lactulose, fucose, rhamnose, melezitose, maltotriose, and raffinose. Preferred sugars include mannitol, lactose, sucrose, sorbitol, trehalose, glucose, and are adjusted to provide osmolality if administered parenterally or to provide wetting of the porous drug matrix or the drug microparticles within the matrix.

(iii) Wetting Agents

25 Wetting agents can be used to facilitate water ingress into the matrix and wetting of the drug particles in order to facilitate dissolution. Representative examples of wetting agents include gelatin, casein, lecithin (phosphatides), gum acacia, cholesterol, tragacanth, stearic acid, benzalkonium chloride, calcium stearate, glycerol monostearate, cetostearyl alcohol, cetomacrogol emulsifying wax, sorbitan esters, polyoxyethylene alkyl ethers (e.g., macrogol ethers such as cetomacrogol 1000), polyoxyethylene castor oil derivatives, polyoxyethylene sorbitan fatty acid esters (e.g., TWEEN<sup>TM</sup>s), polyethylene glycols, polyoxyethylene stearates,

30



colloidal silicon dioxide, phosphates, sodium dodecylsulfate, carboxymethylcellulose calcium, carboxymethylcellulose sodium, methylcellulose, hydroxyethylcellulose, hydroxy propylcellulose, hydroxypropylmethylcellulose phthlate, noncrystalline cellulose, magnesium  
5 aluminum silicate, triethanolamine, polyvinyl alcohol, and polyvinylpyrrolidone (PVP). Tyloxapol (a nonionic liquid polymer of the alkyl aryl polyether alcohol type, also known as superinone or triton) is another useful wetting agent. Most of these wetting agents are known pharmaceutical excipients and are described in detail in the Handbook of  
10 Pharmaceutical Excipients, published jointly by the American Pharmaceutical Association and The Pharmaceutical Society of Great Britain (The Pharmaceutical Press, 1986).

Preferred wetting agents include polyvinylpyrrolidone, polyethylene glycol, tyloxapol, poloxamers such as PLURONIC<sup>TM</sup> F68, F127, and F108,  
15 which are block copolymers of ethylene oxide and propylene oxide, and polyamines such as TETRONIC<sup>TM</sup> 908 (also known as POLOXAMINE<sup>TM</sup> 908), which is a tetrafunctional block copolymer derived from sequential addition of propylene oxide and ethylene oxide to ethylenediamine (available from BASF), dextran, lecithin, dialkylesters of sodium sulfosuccinic acid  
20 such as AEROSOL<sup>TM</sup> OT, which is a dioctyl ester of sodium sulfosuccinic acid (available from American Cyanamid), DUPONOL<sup>TM</sup> P, which is a sodium lauryl sulfate (available from DuPont), TRITON<sup>TM</sup> X-200, which is an alkyl aryl polyether sulfonate (available from Rohm and Haas), TWEEN<sup>TM</sup> 20 and TWEEN<sup>TM</sup> 80, which are polyoxyethylene sorbitan fatty  
25 acid esters (available from ICI Specialty Chemicals), Carbowax 3550 and 934, which are polyethylene glycols (available from Union Carbide), Crodesta F-110, which is a mixture of sucrose stearate and sucrose distearate, and Crodesta SL-40 (both available from Croda Inc.), and SA90HCO, which is  $C_{18}H_{37}CH_2(CON(CH_3)CH_2(CHOH)_4CH_2OH)_2$ .

30 Wetting agents which have been found to be particularly useful include Tetronic 908, the Tweens, Pluronic F-68 and polyvinylpyrrolidone. Other useful wetting agents include decanoyl-N-methylglucamide; n-decyl- $\beta$ -D-glucopyranoside; n-decyl- $\beta$ -D-maltopyranoside; n-dodecyl- $\beta$ -D-

glucopyranoside; n-dodecyl  $\beta$ -D-maltoside; heptanoyl-N-methylglucamide; n-heptyl- $\beta$ -D-glucopyranoside; n-heptyl- $\beta$ -D-thioglucoside; n-hexyl- $\beta$ -D-glucopyranoside; nonanoyl-N-methylglucamide; n-nonyl- $\beta$ -D-glucopyranoside; octanoyl-N-methylglucamide; n-octyl- $\beta$ -D-glucopyranoside; and octyl- $\beta$ -D-thioglucopyranoside. Another preferred wetting agent is p-isononylphenoxypoly(glycidol), also known as Olin-10G or Surfactant 10-G (commercially available as 10G from Olin Chemicals). Two or more wetting agents can be used in combination.

(iv) Tonicity or Osmolality Agents

10 The porous drug matrices may include one or more tonicity agents, such as salts (e.g., as sodium chloride or potassium chloride) or sugars (such as mannitol, dextrose, sucrose, or trehalose) to adjust a hypotonic solution of a drug to isotonic so that the drug, when in solution, is physiologically compatible with the cells of the body tissue of the patient. The type and  
15 amount of tonicity agent can be selected by one of skill in the art using known techniques.

(v) Pegylated Excipients

In one embodiment, the matrix further includes a pegylated excipient. Such pegylated excipients include, but are not limited to, pegylated  
20 phospholipids, pegylated proteins, pegylated peptides, pegylated sugars, pegylated polysaccharides, pegylated block copolymers with one of the blocks being PEG, and pegylated hydrophobic compounds such as pegylated cholesterol. The pegylated excipient beneficially envelops or shields the drug from macrophage uptake, which prolongs its half-life or enhances  
25 bioavailability of the drug.

Representative examples of pegylated phospholipids include 1,2-diacyl-sn-glycero-3-phosphoethanolamine-N-[Poly(ethylene glycol) 2000] ("PEG 2000 PE") and 1,2-diacyl-sn-glycero-3-phosphoethanolamine-N-[Poly(ethylene glycol) 5000] ("PEG 5000 PE"), where the acyl group is  
30 selected, for example, from dimyristoyl, dipalmitoyl, distearoyl, dioleoyl, and 1-palmitoyl-2-oleoyl.

Other polyalkyleneoxides can be used in place of the polyethylene.

## **II. Volatile Solvents**

The choice of solvent depends on the drug. In a preferred embodiment, the solvent is an organic solvent that is volatile, has a relatively low boiling point, or can be removed under vacuum, and which is acceptable for administration to humans in trace amounts. Representative solvents include acetic acid, acetaldehyde dimethyl acetal, acetone, acetonitrile, chloroform, chlorofluorocarbons, dichloromethane, dipropyl ether, diisopropyl ether, N,N-dimethylformamide (DMF), formamide, dimethyl sulfoxide (DMSO), dioxane, ethanol, ethyl acetate, ethyl formate, ethyl vinyl ether, methyl ethyl ketone (MEK), glycerol, heptane, hexane, isopropanol, methanol, isopropanol, butanol, triethylamine, nitromethane, octane, pentane, tetrahydrofuran (THF), toluene, 1,1,1-trichloroethane, 1,1,2-trichloroethylene, water, xylene, and combinations thereof. In general, the drug is dissolved in the volatile solvent to form a drug solution having a concentration of between 0.01 and 80% weight to volume (w/v), more preferably between 0.025 and 30% (w/v).

Aqueous solvents or mixtures of aqueous and organic solvents, such as water-alcohol mixtures, can be used to dissolve the drug.

## **III. Pore Forming Agents**

Pore forming agents are volatile materials that are used during the process to create porosity in the resultant matrix. The pore forming agent can be a volatilizable solid or volatilizable liquid.

### **1. Liquid Pore Forming Agent**

The liquid pore forming agent must be immiscible with the drug solvent and volatilizable under processing conditions compatible with the drug. To effect pore formation, the pore forming agent first is emulsified with the drug solvent. Then, the emulsion is further processed to remove the drug solvent and the pore forming agent simultaneously or sequentially using evaporation, vacuum drying, spray drying, fluid bed drying, lyophilization, or a combination of these techniques.

The selection of liquid pore forming agents will depend on the drug solvent. Representative liquid pore forming agents include water; dichloromethane; alcohols such as ethanol, methanol, or isopropanol;

acetone; ethyl acetate; ethyl formate; dimethylsulfoxide; acetonitrile; toluene; xylene; dimethylformamide; ethers such as THF, diethyl ether, or dioxane; triethylamine; formamide; acetic acid; methyl ethyl ketone; pyridine; hexane; pentane; furan; water; and cyclohexane.

- 5           The liquid pore forming agent is used in an amount that is between 1 and 50% (v/v), preferably between 5 and 25% (v/v), of the drug solvent emulsion.

## 2. Solid Pore Forming Agent

- 10           The solid pore forming agent must be volatilizable under processing conditions which do not harm the drug compositions. The solid pore forming agent can be (i) dissolved in the drug solution, (ii) dissolved in a solvent which is not miscible with the drug solvent to form a solution which is then emulsified with the drug solution, or (iii) added as solid particulates to the drug solution. The solution, emulsion, or suspension of the pore  
15           forming agent in the drug solution then is further processed to remove the drug solvent, the pore forming agent, and, if appropriate, the solvent for the pore forming agent simultaneously or sequentially using evaporation, spray drying, fluid bed drying, lyophilization, vacuum drying, or a combination of these techniques.

- 20           In a preferred embodiment, the solid pore forming agent is a volatile salt, such as salts of volatile bases combined with volatile acids. Volatile salts are materials that can transform from a solid or liquid to a gaseous state using added heat and/or vacuum. Examples of volatile bases include ammonia, methylamine, ethylamine, dimethylamine, diethylamine,  
25           methylethylamine, trimethylamine, triethylamine, and pyridine. Examples of volatile acids include carbonic acid, hydrochloric acid, hydrobromic acid, hydroiodic acid, formic acid, acetic acid, propionic acid, butyric acid, and benzoic acid. Preferred volatile salts include ammonium bicarbonate, ammonium acetate, ammonium chloride, ammonium benzoate and mixtures  
30           thereof.

Other examples of solid pore forming agents include iodine, phenol, benzoic acid (as acid not as salt), and naphthalene.

The solid pore forming agent is used in an amount between 0.5 and

1000% (w/w), preferably between 10 and 600% (w/w), and more preferably between 1 and 100% (w/w), of the drug.

#### IV. Method of Making the Porous Drug Matrix

The porous drug matrices preferably are made by (i) dissolving a  
5 drug, preferably one having low aqueous solubility, in a volatile solvent to form a drug solution, (ii) combining at least one pore forming agent with the drug solution to form an emulsion, suspension, or second solution, and (iii) removing the volatile solvent and pore forming agent from the emulsion, suspension, or second solution. In a preferred embodiment, spray drying,  
10 optionally followed by lyophilization or vacuum drying, is used to remove the solvents and the pore forming agent. The removal of the pore forming agent can be conducted simultaneously with or following removal of enough solvent to solidify the droplets. Production can be carried out using continuous, batch, or semi-continuous processes.

15 First, the selected drug is dissolved in an appropriate solvent. The concentration of the drug in the resulting drug solution typically is between about 0.01 and 80% (w/v), preferably between about 0.025 and 30% (w/v).

Next, the drug solution is combined, typically under mixing conditions, with the pore forming agent or solution thereof. If a liquid pore  
20 forming agent is used, it is first emulsified with the drug solution to form droplets of pore forming agent dispersed throughout the drug solution. If a solid pore forming agent is used, it is dissolved either directly in the drug solution to form a solution of drug/pore forming agent, or it is first dissolved in a second solvent. If the second solvent is immiscible with the drug  
25 solvent, the solution of the pore forming agent is emulsified with the drug solution to form droplets of the pore forming agent solution dispersed throughout the drug solution. If the second solvent is miscible with the drug solution, the two solutions are mixed to form a single drug solution. A solid pore forming agent alternatively can be added directly to the drug solution as  
30 solid particulates, preferably between about 100 nm and 10  $\mu$ m in size, to form a suspension of pore forming agent in the drug solution. Subsequently, the solid pore forming agent particle size can be reduced by further processing the resulting suspension, for example, using homogenization or

sonication techniques known in the art.

Then, the solution, emulsion, or suspension is further processed to remove the drug solvent and the pore forming agent simultaneously or sequentially, using evaporation, spray drying, fluid bed drying, lyophilization, vacuum drying, or a combination of these techniques. In a preferred embodiment, the solution, emulsion, or suspension is spray-dried. As used herein, "spray dry" means to atomize the solution, emulsion, or suspension to form a fine mist of droplets (of drug solution having solid or liquid pore forming agent dispersed throughout), which immediately enter a drying chamber (e.g., a vessel, tank, tubing, or coil) where they contact a drying gas. The solvent and pore forming agents evaporate from the droplets into the drying gas to solidify the droplets, simultaneously forming pores throughout the solid. The solid (typically in a powder, particulate form) then is separated from the drying gas and collected.

The temperature of the inlet and outlet ports of the drying chamber, as well as the flow rates of the feed solution, atomization gas, and drying gas, can be controlled to produce the desired products. In a particularly preferred embodiment, the spray drying methods described in U.S. Patent No. 5,853,698 to Straub et al. are adapted to make the drug matrices.

The drug present in the solids or powder produced may be in a crystalline or an amorphous state, or may be mixture of such states. The state generally depends on how the droplets are dried and the excipients present.

#### Emulsion Stabilization

In embodiments in which at least one pore forming agent is combined with the drug solution to form an emulsion, a surfactant or emulsifying agent can be added to enhance the stability of the emulsion. A variety of surfactants may be incorporated in this process, preferably to an amount between 0.1 and 5% by weight. Exemplary emulsifiers or surfactants which may be used include most physiologically acceptable emulsifiers, for instance egg lecithin or soya bean lecithin, or synthetic lecithins such as saturated synthetic lecithins, for example, dimyristoyl phosphatidyl choline, dipalmitoyl phosphatidyl choline or distearoyl phosphatidyl choline or

unsaturated synthetic lecithins, such as dioleoyl phosphatidyl choline or  
dilinoleoyl phosphatidyl choline. Other hydrophobic or amphipathic  
compounds can be used in place of the phospholipid, for example,  
cholesterol. Emulsifiers also include surfactants such as free fatty acids,  
5 esters of fatty acids with polyoxyalkylene compounds like polyoxpropylene  
glycol and polyoxyethylene glycol; ethers of fatty alcohols with  
polyoxyalkylene glycols; esters of fatty acids with polyoxyalkylated  
sorbitan; soaps; glycerol-polyalkylene stearate; glycerol-polyoxyethylene  
ricinoleate; homo- and co-polymers of polyalkylene glycols; polyethoxylated  
10 soya-oil and castor oil as well as hydrogenated derivatives; ethers and esters  
of sucrose or other carbohydrates with fatty acids, fatty alcohols, these being  
optionally polyoxyalkylated; mono-, di- and tri-glycerides of saturated or  
unsaturated fatty acids, glycerides of soya-oil and sucrose.

Other emulsifiers include natural and synthetic forms of bile salts or  
15 bile acids, both conjugated with amino acids and unconjugated such as  
taurodeoxycholate and cholic acid.

#### **V. Porous Drug Matrix Applications**

The porous drug matrices described herein are useful in formulations  
for administration to a patient in need of the drug. As used herein, "patient"  
20 refers to animals, including mammals, preferably humans. The formulations  
deliver a therapeutically or prophylactically effective amount of the drug to  
the patient.

In a preferred embodiment, paclitaxel or docetaxel matrices are used  
in formulations for administration to a patient in need thereof.  
25 Pharmaceutical compositions of these drugs preferably are provided without  
CREMOPHOR<sup>TM</sup>, Polysorbate 80, or other solubilizing agents, for bolus  
administration. The compositions are porous dry powders which upon the  
addition of an aqueous medium form a suspension of nanoparticles and  
microparticles of the drug.

30 The porous matrices, or formulations thereof, are suitable for  
administration of drug by a variety of routes, for example, parenteral,  
mucosal, oral, topical/transdermal administration, for local, regional, or  
systemic effect. Examples of parenteral routes include intravenous,

intraarterial, intracardiac, intrathecal, intraosseous, intraarticular, intrasynovial, intracutaneous, subcutaneous, and intramuscular administration. Examples of mucosal routes include pulmonary (intrarespiratory), buccal, sublingual, intranasal, rectal, and vaginal administration. The porous matrices also can be formulated for intraocular, conjunctival, aural, urethral, intracranial, intralesional, and intratumoral administration.

In a preferred embodiment, the drug matrix is in the form of powder, which can be reconstituted with an aqueous medium, such as physiological saline, and administered parenterally, such as intramuscularly, subcutaneously, or intravenously. An advantage of the formulations described herein is that they can be used to convert drugs which must be infused (e.g., to avoid precipitation of the drug following bolus injection) to a bolus formulation, avoiding unacceptable precipitation of drug *in vivo* or for local delivery.

Alternatively, the matrix can be further processed using standard techniques into tablets or capsules for oral administration, into rectal suppositories, into a dry powder inhaler for pulmonary administration, or mixed/processed into a cream or ointment for topical administration. These standard techniques are described, for example, in Ansel, et al., "*Pharmaceutical Dosage Forms and Drug Delivery Systems*," 6<sup>th</sup> Ed., (Williams & Wilkins 1995).

The present invention will be further understood with reference to the following non-limiting examples.

## Overview

Examples 1-10 demonstrate production of porous drug matrices using different pore forming agents, different drugs, and different solvents. Examples 1-8 use emulsion formulations to produce the matrices, whereas Examples 9 and 10 use solution formulations to produce the matrices. Examples 11-13 describe the analyses which were used to characterize the porous drug matrices produced in Examples 1-10. These characteristics include density, drug integrity, and dissolution properties.

Example 14 describes particle size analysis and surface area analysis



of low water solubility drug particles incorporated into the porous drug matrices. Examples 15-17 describe experiments demonstrating the increased internal surface area of porous drug matrices produced with pore forming agents. Examples 18-21 describe experiments demonstrating the advantage or need to include a wetting agent as a component of the porous drug matrices. Example 22 describes an experiment demonstrating the administration of porous drug matrices as an intravenous bolus. Examples 23 and 24 describe the production of porous drug matrices produced with pore forming agents and pegylated phospholipids.

#### 10        Materials and Equipment

The following materials and equipment were used in the examples. PEG 3350, PEG 8000, polyvinylpyrrolidone K-15, nifedipine, naproxen, prednisone, SPAN<sup>TM</sup> 40, lecithin, TWEEN<sup>TM</sup> 80, PLURONIC<sup>TM</sup> F127, ammonium chloride, ammonium bicarbonate, and ammonium acetate were obtained from Spectrum Chemicals (Gardena, CA). Griseofulvin was obtained from Aldrich Chemicals (Milwaukee, WI). Paclitaxel was obtained from Hauser (Boulder, CO). 1,2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine-N-[Poly(ethylene glycol)-5000] (PEG 5000 PE) and 1,2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine-N-[Poly(ethylene glycol)-2000] (PEG 2000 PE) were obtained from Avanti Polar Lipids Inc. (Alabaster, AL). Methylene chloride was obtained from EM Science (Gibbstown, NJ). All emulsions were produced using a Virtis IQ<sup>2</sup> homogenizer (Virtis, Gardiner, NY). Formulations were spray dried on a benchtop spray dryer using an air atomizing nozzle.

#### 25        **Example 1:    Production of a Porous Prednisone Matrix Using                          Ammonium Bicarbonate as a Pore Forming Agent with                          SPAN<sup>TM</sup> 40 and PEG 8000 as Wetting Agents**

5.46 g of PEG 8000, 0.545 g of prednisone, and 0.055 g of SPAN<sup>TM</sup> 40 were dissolved in 182 mL of methylene chloride. An aqueous solution was prepared by dissolving 3.27 g of ammonium bicarbonate in 18.2 mL of deionized (DI) water. The aqueous solution was added to the organic solution (phase ratio 1:10) and homogenized for 5 minutes at 16,000 RPM. The resulting emulsion was spray dried on a benchtop spray dryer using an

air-atomizing nozzle and nitrogen as the drying gas. Spray drying process conditions were 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 36 °C outlet temperature.

**Example 2: Production of a Porous Prednisone Matrix Using**

5                   **Ammonium Bicarbonate as a Pore Forming Agent with  
PEG 8000, TWEEN™ 80, Lecithin as Wetting Agents**

5.46 g of PEG 8000, 0.545 g of prednisone, 0.003 g of TWEEN™  
80, and 0.003 g of lecithin were dissolved in 182 mL of methylene chloride.  
An aqueous solution was prepared as described in Example 1. The aqueous  
10 solution was added to the organic solution (phase ratio 1:10) and  
homogenized for 15 minutes as described in Example 1. The resulting  
emulsion was spray dried as described in Example 1 using process  
conditions of 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 35  
°C outlet temperature.

15   **Example 3: Production of a Porous Prednisone Matrix Using  
Ammonium Acetate as a Pore Forming Agent, and PEG  
8000, TWEEN™ 80, and Lecithin as Wetting Agents**

A prednisone-loaded organic solution was prepared as described in  
Example 2. An aqueous solution was prepared by dissolving 3.27 g of  
20 ammonium acetate in 18.2 mL of DI water. The aqueous and organic  
solutions were homogenized and spray dried as described in Example 2.

**Example 4: Production of a Porous Prednisone Matrix Using  
Ammonium Chloride as a Pore Forming Agent, and PEG  
8000, TWEEN™ 80, and Lecithin as Wetting Agents**

25                   A prednisone-loaded organic solution was prepared as described in  
Example 2. An aqueous solution was prepared by dissolving 3.27 g of  
ammonium chloride in 18.2 mL of DI water. The aqueous and organic  
solutions were homogenized as described in Example 1. The resulting  
emulsion was spray dried as described in Example 2.

30   **Example 5: Production of a Porous Griseofulvin Matrix Using  
Ammonium Bicarbonate as a Pore Forming Agent, and  
PEG 3350, TWEEN™ 80, and Lecithin as Wetting Agents**

9.09 g of PEG 3350, 4.55 g of griseofulvin, 0.01 g of TWEEN™ 80,

and 0.01 g of lecithin were dissolved in 182 mL of methylene chloride. An aqueous solution was prepared by dissolving 3.27 g of ammonium bicarbonate and 1.09 g of PEG 3350 in 18.2 mL of DI water. The aqueous and organic solutions were homogenized as described in Example 1. The resulting emulsion was spray dried as described in Example 1 using process conditions of 20 ml/min solution flow rate, 80 kg/hr drying gas rate, and 12 °C outlet temperature.

**Example 6: Production of a Porous Nifedipine Matrix Using Ammonium Bicarbonate as a Pore Forming Agent, and PEG 3350 and Lecithin as Wetting Agents**

9.09 g of PEG 3350, 2.27 g of nifedipine, and 0.009 g of lecithin were dissolved in 182 mL of methylene chloride. An aqueous solution was prepared by dissolving 3.27 g of ammonium bicarbonate in 18.2 mL of DI water. The aqueous and organic solutions were homogenized in described in Example 1. The resulting emulsion was spray dried as described in Example 1 using process conditions of 20 ml/min solution flow rate, 60 kg/hr drying gas rate, and 20 °C outlet temperature.

**Example 7: Production of a Porous Naproxen Matrix Using Ammonium Chloride as a Pore Forming Agent, and PEG 3350 and Lecithin as Wetting Agents**

A naproxen-loaded organic solution was prepared by dissolving 10.91 g of PEG 3350, 2.73 g of naproxen, and 0.109 g of lecithin in 182 mL of methylene chloride. An aqueous solution was prepared as described in Example 4. The aqueous and organic solutions were homogenized as described in Example 1, and the resulting emulsion was spray dried using process conditions of 20 ml/min solution flow rate, 100 kg/hr drying gas rate, and 20 °C outlet temperature.

**Example 8: Production of a Porous Paclitaxel Matrix Using Ammonium Bicarbonate as a Pore Forming Agent, and PEG 3350 and Lecithin as Wetting Agents**

A paclitaxel-loaded organic solution was prepared by dissolving 3.0 g of paclitaxel, 15.0 g of PEG 3350, and 15.7 mg of lecithin in 100 mL of methylene chloride. An aqueous solution composed of 1.8 g of ammonium

bicarbonate and 0.6 g of PEG 3350 in 10 mL of DI water was added to the organic solution (phase ratio 1:10). The mixture was homogenized for 5 minutes at 16,000 RPM. The resulting emulsion was spray dried using process conditions of 10 mL/min solution flow rate, 60 kg/hr drying gas rate, and 25 °C outlet temperature.

**Example 9: Production of a Porous Nifedipine Matrix Using Ammonium Bicarbonate as a Pore Forming Agent, PEG 3350 and TWEEN™ 80 as Wetting Agents, Polyvinylpyrrolidone as a Bulking Agent, and Ethanol as a Solvent**

A nifedipine-loaded organic solution was prepared by dissolving 0.76 g of nifedipine, 0.28 g of PEG 3350, and 2.72 g of polyvinylpyrrolidone K-15 in 170 mL of ethanol. An aqueous solution composed of 1.62 g of ammonium bicarbonate and 3 mg of TWEEN™ 80 in 30 mL of DI water was added to the ethanol solution and mixed. The resulting solution was spray dried using process conditions of 20 mL/min solution flow rate, 100 kg/hr drying gas rate, and 36 °C outlet temperature.

**Example 10: Production of a Porous Nifedipine Matrix Using Ammonium Bicarbonate as a Pore Forming Agent, PEG 3350 and PLURONIC™ F127 as Wetting Agents, Polyvinylpyrrolidone as a Bulking Agent, and Ethanol as a Solvent**

A nifedipine-loaded organic solution was prepared by dissolving 0.76 g of nifedipine, 0.28 g of PEG 3350, and 2.72 g of polyvinylpyrrolidone K-15 in 170 mL of ethanol. An aqueous solution composed of 1.62 g of ammonium bicarbonate and 3 mg of PLURONIC™ F127 in 30 mL of DI water was added to the ethanol solution and mixed. The resulting solution was spray dried using process conditions of 20 mL/min solution flow rate, 100 kg/hr drying gas rate, and 36 °C outlet temperature.

**Example 11: In Vitro Dissolution of Porous Drug Matrices**

The *in vitro* dissolution rates of the powders produced in Examples 1-10 were compared to the dissolution rates of the bulk drug of interest.

### Analytical Method

All dissolution studies were conducted in PBS (phosphate buffered saline) at room temperature in a glass beaker using overhead mixing. The mixer used was an IKARW16 Basic Mixer with a R1342 impeller shaft  
5 running at stirring rate 5. Samples were removed via pipet, filtered through 0.22 micron CA syringe filter, and then analyzed. UV-vis spectroscopy was conducted on an Hewlett Packard Model 8453. Dissolution curves are presented as percent of complete dissolution.

For griseofulvin, PBS (600 mL) was added to an appropriate amount  
10 of material being tested to contain 2.4 mg of griseofulvin. UV analysis was performed at 291 nm.

For naproxen, PBS (100 mL) was added to an appropriate amount of material being tested to contain 100 mg of naproxen. All vessels containing naproxen as a solid or as a solution were protected from light. UV analysis  
15 was performed at 332 nm.

For nifedipine, PBS (600 mL) was added to an appropriate amount of material being tested to contain 2.4 mg of nifedipine. All vessels containing nifedipine as a solid or in solution were protected from light. UV analysis was performed at 237 nm.

20 For prednisone, PBS (250 mL) was added to an appropriate amount of material being tested to contain 5 mg of prednisone. UV analysis was performed at 244 nm.

For paclitaxel, studies were conducted in PBS containing 0.08% TWEEN<sup>TM</sup> 80 (T80/PBS). T80/PBS (10 mL) was added to an appropriate  
25 amount of material being tested to contain 5 mg of paclitaxel in a 15 mL polypropylene conical tube, and the suspension was vortexed for 3-4 minutes. The suspension (0.25 mL) was then added to 250 mL of T80/PBS in a 600 mL glass beaker for dissolution analysis. HPLC analysis was performed directly on the filtered aqueous solutions using the paclitaxel  
30 HPLC method described in Example 13.

### Results

The *in vitro* dissolution rates of the porous drug matrices produced in examples 1-10 are provided in Figures 1-6. The *in vitro* dissolution of the

porous drug matrices are compared to the bulk drug of interest. In all cases, the time for 80% dissolution of the porous drug matrices is 4-50 times shorter than the time for 80% of the bulk drug to dissolve. The rate of dissolution which is approximated as the slope of the curve is 10 to 1400 times greater for the porous drug matrices of Examples 1-10 as compared to the specific bulk drug of interest.

#### Example 12: Density of Porous Drug Matrices

The densities of the dry powder produced in Examples 1-7 are summarized in Table 1. Density was measured using Transaxial Pressure ("TAP") with a Micromeritics GeoPyc 1360 using a consolidation force of 8 Newtons. The matrices are less dense than the starting bulk drug in all cases, indicating that the porous drug matrices are more porous than the commercially available bulk drug.

Table 1: Particle Density Analysis

Material	Density (g/mL)
Prednisone Bulk	0.68
Example 1	0.48
Example 2	0.55
Example 3	0.51
Example 4	0.49
Griseofulvin Bulk	0.80
Example 5	0.55
Nifedipine Bulk	1.01
Example 6	0.56
Naproxen Bulk	0.69
Example 7	0.58

#### Example 13: Integrity of the Drug in Porous Drug Matrices

##### Analytical Method

Drug integrity post processing was assessed by High Pressure Liquid Chromatography ("HPLC") (Hewlett Packard Series 1100 HPLC). USP chromatography conditions were used for prednisone, naproxen, nifedipine, and griseofulvin. Vessels and vials containing naproxen or nifedipine solutions were protected from light. For paclitaxel, the chromatographic conditions included a Nucleosil column (5  $\mu$ m, C18, 100A, 250 x 4.6 mm), a mobile phase of 2 mM H<sub>3</sub>PO<sub>4</sub>/Acetonitrile (2:3) at a flow rate of 1.5 mL/min, UV detection at 227 nm, and a run time of 25 min.

### Results

The integrities of the drugs following the processing in Examples 1-9 are shown in Table 2 as purities. The process of forming the drug into porous matrices does not appear to alter the purity of the drug.

5 **Table 2: Drug Integrity Analysis**

Material	Purity (%)
Prednisone Powder	100
Example 1	99.8
Example 2	99.8
Example 3	99.8
Example 4	99.8
Griseofulvin Bulk	95.7
Example 5	95.7
Nifedipine Bulk	100
Example 6	100
Example 9	100
Example 10	100
Naproxen Bulk	100
Example 7	100
Paclitaxel Bulk	100
Example 8	100

### **Example 14: Particle Size Analysis and Surface Area Analysis of Drug Particles in Wetted Porous Drug Matrices**

#### Analytical Methods

10 Particle size analysis was performed using the Coulter Multisizer II with a 50 micron aperture using siphon mode. Electrolyte was pre-saturated with the drug of interest, and filtered through a 0.22 micron filter prior to addition of lots for analysis to ensure that no portion of the drug within the lot would dissolve during the analysis.

#### 15 Results

The mean particle size and total surface area of the drug particles generated when the porous drug matrices produced in Examples 1-7 were reconstituted in aqueous media are summarized in Table 3.

**Table 3: Particle Size and Surface Area Analysis**

<b>Material</b>	<b>Size (microns)</b>	<b>Surface Area (m<sup>2</sup>/mL of microparticles)</b>
Prednisone Powder	2.07	1.43
Example 1	1.58	1.66
Example 2	1.39	2.53
Example 3	1.39	3.02
Example 4	1.24	3.36
Griseofulvin Bulk	2.42	0.88
Example 5	2.16	1.28
Nifedipine Bulk	2.64	0.57
Example 6	1.78	1.98
Naproxen Bulk	2.89	0.66
Example 7	1.34	2.79

In all cases, the particle size of the drug particles which resulted from wetting of the porous drug matrices was reduced relative to the starting bulk material by 10 to 54%, and the total surface area of the particles was increased relative to the starting bulk drug by approximately 16-320%.

**Example 15: Nifedipine Drug Matrices Containing a Wetting Agent**

**Produced With and Without a Pore Forming Agent**

A nifedipine/PEG solution was prepared by dissolving 2.0 g of nifedipine, 8.0 g of PEG 3350, and 8 mg of lecithin in 200 mL of methylene chloride (Example 15A). A second identical nifedipine-loaded organic solution was prepared. An aqueous solution composed of 1.8 g of ammonium bicarbonate in 20 mL of DI water was added to the first nifedipine organic solution (phase ratio 1:10). The mixture was homogenized for 5 minutes at 16,000 RPM. The nifedipine solution (Example 15A) and the nifedipine emulsion (Example 15B) were separately spray dried using process conditions of 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 21 °C outlet temperature.

**Example 16: Griseofulvin Drug Matrices Containing a Wetting Agent**

**Produced With and Without a Pore Forming Agent**

A griseofulvin/PEG solution was prepared by dissolving 5.0 g of griseofulvin, 11.2 g of PEG 3350, 11 mg of TWEEN<sup>TM</sup> 80, and 11 mg of lecithin in 200 mL of methylene chloride (Example 16A). A second identical griseofulvin-loaded organic solution was prepared. An aqueous



solution composed of 1.8 g of ammonium bicarbonate in 20 mL of DI water was added to the first organic solution (phase ratio 1:10). The mixture was homogenized for 5 minutes at 16,000 RPM. The griseofulvin solution (Example 16A) and griseofulvin emulsion (Example 16B) were spray dried on a benchtop spray dryer using process conditions of 20 mL/min solution flow rate, 80 kg/hr drying gas rate, and 13 °C outlet temperature.

**Example 17: Internal Surface Area of Porous Drug Matrices  
Containing a Wetting Agent and Produced With  
and Without a Pore Forming Agent**

The internal surface areas of the drug matrices produced in Examples 15 and 16 were assessed by Krypton BET. BET specific surface area analysis was performed using multi-point surface area analysis with krypton as the gas. Samples were outgassed to 20 micron vacuum at 20 °C prior to analysis. The results, shown in Table 4, illustrate that the use of the pore forming agent led to an increase of between 2.3 and 3.5 fold in the internal surface area of the resultant drug matrix.

**Table 4: Internal Surface Area of Drug Matrices**

Matrix (Example No.)	Surface Area (m <sup>2</sup> /g matrix)
Nifedipine with wetting agent (15A)	0.40
Nifedipine with wetting agent and Ammonium Bicarbonate (15B)	1.4
Griseofulvin with wetting agent (16A)	0.41
Griseofulvin with wetting agent and Ammonium Bicarbonate (16B)	0.95

**Example 18: Nifedipine Drug Matrix Produced Without a Pore  
Forming Agent or Wetting Agent**

A 5% nifedipine solution was prepared by dissolving 10.0 g of nifedipine in 200 mL of methylene chloride. The solution was spray dried on a benchtop spray dryer using the following conditions: 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 22°C outlet temperature.

**Example 19: Griseofulvin Drug Matrix Produced Without a Pore  
Forming Agent or Wetting Agent**

An 8.1% griseofulvin solution was prepared by dissolving 16.2 g of griseofulvin in 200 mL of methylene chloride. The solution was spray dried

on a benchtop spray dryer using process conditions of 20 mL/min solution flow rate, 80 kg/hr drying gas rate, and 13 °C outlet temperature.

**Example 20: *In Vitro* Dissolution of Nifedipine Drug Matrices Produced With / Without Pore Forming Agent and Wetting Agent**

5        The *in vitro* dissolution rates of the nifedipine matrices produced in Examples 15 and 18 are shown in Figure 6. The *in vitro* dissolution of the drug matrices produced with either wetting agent or wetting agent and pore forming agent have increased dissolution rates as compared to the drug matrix produced with the drug alone. The matrix produced with both the  
10        wetting agent and the pore forming agent has the greatest dissolution rate.

**Example 21: *In Vitro* Dissolution of Griseofulvin Drug Matrices Produced With / Without Pore Forming Agent and Wetting Agent**

15        The *in vitro* dissolution rates of the griseofulvin matrices produced in examples 16 and 19 are provided in Figure 7. The *in vitro* dissolution of the drug matrices produced with either wetting agent or wetting agent and pore forming agent have increased dissolution rates as compared to the drug matrix produced with the drug alone. The matrix produced with both the wetting agent and the pore forming agent has the greatest dissolution rate.

20        **Example 22: Administration of Porous Drug Matrices as an Intravenous Bolus to Dogs**

25        A nifedipine-loaded organic solution was prepared by dissolving 9.09 g of PEG 3350, 2.27 g of nifedipine, and 0.009 g of lecithin in 182 mL of methylene chloride. An aqueous solution was prepared by dissolving 3.27 g of ammonium bicarbonate and 0.91 g of PEG 3350 in 18.2 mL of deionized water at room temperature. The aqueous and organic solutions were homogenized as described in Example 1, and the resulting emulsion was spray dried using process conditions of 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 20 °C outlet temperature.

30        A suspension of the porous nifedipine drug matrix was prepared in 5% dextrose solution at a concentration of 2.5 mg/mL. The suspension (2 mL) was administered as a bolus to four beagle dogs, which weighed 8-10 kg. Blood samples were taken at time-points ranging from 1 minute to 24

hours. The samples were processed into plasma, were stored frozen, and were protected from light until analysis via liquid chromatography-mass spectrometry. All animals tolerated the suspension administered as a bolus. The average plasma levels of the intravenously administered suspension is shown in Figure 8.

**Example 23: Production of a Porous Nifedipine Matrix Using a Pegylated Phospholipid, 1, 2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine-N-[Poly(ethylene glycol)-5000]**

A nifedipine-loaded organic solution was prepared by dissolving 2.0 g of nifedipine, 30.0 g of PEG 3350, 4 mg of lecithin, and 4 mg of 1, 2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine-N-[Poly(ethylene glycol)-5000] (PEG 5000 PE) in 202 mL of methylene chloride. An aqueous solution of 1.8 g of ammonium bicarbonate in 20 mL of DI water was added to the organic solution (phase ratio 1:10). The mixture was homogenized for 5 minutes at 16,000 RPM. The resulting emulsion was spray dried using process conditions of 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 21 °C outlet temperature.

**Example 24: Production of a Porous Nifedipine Matrix Using a Pegylated Phospholipid, 1, 2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine-N-[Poly(ethylene glycol)-2000]**

A nifedipine-loaded organic solution was prepared by dissolving 2.0 g of nifedipine, 30.0 g of PEG 3350, 4 mg of lecithin, and 4 mg of 1, 2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine-N-[Poly(ethylene glycol)-2000] (PEG 2000 PE) in 202 mL of methylene chloride. An aqueous solution composed of 1.8 g of ammonium bicarbonate in 20 ml of DI water was added to the organic solution (phase ratio 1:10). The mixture was homogenized for 5 minutes at 16,000 RPM. The resulting emulsion was spray dried using process conditions of 20 mL/min solution flow rate, 60 kg/hr drying gas rate, and 21 °C outlet temperature.

Modifications and variations of the present invention will be obvious to those of skill in the art from the foregoing detailed description. Such modifications and variations are intended to come within the scope of the following claims.

We claim:

1. A method for making a porous matrix of drug comprising
  - (a) dissolving a drug in a volatile solvent to form a drug solution,
  - (b) combining at least one pore forming agent with the drug solution to form an emulsion, suspension, or second solution, and
  - (c) removing the volatile solvent and pore forming agent from the emulsion, suspension, or second solution to yield the porous matrix of drug.
2. The method of claim 1 wherein the drug has low aqueous solubility.
3. The method of claim 1 further comprising incorporating at least one wetting agent into the emulsion, suspension, or second solution.
4. The method of claim 1 wherein step (c) is conducted using a process selected from spray drying, evaporation, fluid bed drying, lyophilization, vacuum drying, or a combination thereof.
5. The method of claim 1 wherein the drug solution or pore forming agent further comprises an excipient selected from the group consisting of hydrophilic polymers, sugars, pegylated excipients, and tonicity agents.
6. The method of claim 1 wherein the pore forming agent is a volatile salt.
7. The method of claim 6 wherein the volatile salt is selected from the group consisting of ammonium bicarbonate, ammonium acetate, ammonium chloride, ammonium benzoate, and mixtures thereof.
8. The method of claim 3 which yields a porous matrix formed of a wetting agent and microparticles of a drug, wherein the microparticles have a mean diameter between about 0.01 and 5  $\mu\text{m}$  and a total surface area greater than about 0.5  $\text{m}^2/\text{mL}$ , and wherein the porous matrix is in a dry powder form.
9. A pharmaceutical composition comprising a porous matrix formed of a wetting agent and microparticles of a drug,
  - wherein the microparticles have a mean diameter between about 0.01 and 5  $\mu\text{m}$  and a total surface area greater than about 0.5  $\text{m}^2/\text{mL}$ , and
  - wherein the dry porous matrix is in a dry powder form.
10. The composition of claim 9 wherein the porous matrix is in a dry

powder form having a TAP density less than or equal to 1.0 g/mL.

11. The composition of claim 9 wherein the drug is a low aqueous solubility drug.

12. The composition of claim 11 wherein the matrix upon contact with an aqueous medium yields microparticles having a mean diameter between about 0.1 and 5  $\mu\text{m}$  and a total surface area greater than about 0.5  $\text{m}^2/\text{mL}$ , and wherein the dry porous matrix is in a dry powder form having a TAP density less than or equal to 1.0 g/mL.

13. The composition of claim 9 wherein the drug is a taxane.

14. The composition of claim 13 wherein the taxane is paclitaxel or docetaxel.

15. The composition of claim 11 wherein the drug is selected from the group consisting of albuterol, adapalene, budesonide, doxazosin mesylate, mometasone furoate, ursodiol, amphotericin, enalapril maleate, felodipine, nefazodone hydrochloride, valrubicin, albendazole, estrogens conjugated, medroxyprogesterone acetate, nicardipine hydrochloride, zolpidem tartrate, amlodipine besylate, ethinyl estradiol, omeprazole, rubitecan, amlodipine besylate/ benazepril hydrochloride, etodolac, paroxetine hydrochloride, atovaquone, felodipine, podofilox, paricalcitol, betamethasone dipropionate, fentanyl, pramipexole dihydrochloride, Vitamin D<sub>3</sub> and related analogues, finasteride, quetiapine fumarate, alprostadil, candesartan, cilexetil, fluconazole, ritonavir, busulfan, carbamazepine, flumazenil, risperidone, carbamazepine, carbidopa/levodopa, ganciclovir, saquinavir, amprenavir, carboplatin, glyburide, sertraline hydrochloride, rofecoxib carvedilol, halobetasolpropionate, sildenafil citrate, celecoxib, chlorthalidone, imiquimod, simvastatin, citalopram, ciprofloxacin, irinotecan hydrochloride, sparfloxacin, efavirenz, cisapride monohydrate, lansoprazole, tamsulosin hydrochloride, mofafinil, azithromycin, clarithromycin, letrozole, terbinafine hydrochloride, rosiglitazone maleate, diclofenac sodium, lomefloxacin hydrochloride, tirofiban hydrochloride, telmisartan, diazepam, loratadine, toremifene citrate, thalidomide, dinoprostone, mefloquine hydrochloride, trandolapril, mitoxantrone hydrochloride, tretinoin, etodolac, triamcinolone acetate, estradiol, ursodiol, nelfinavir mesylate, indinavir,

beclomethasone dipropionate, oxaprozin, flutamide, famotidine, nifedipine, prednisone, cefuroxime, lorazepam, digoxin, lovastatin, griseofulvin, naproxen, ibuprofen, isotretinoin, tamoxifen citrate, nimodipine, amiodarone, and alprazolam.

16. The composition of claim 9 wherein the drug is water soluble.

17. The composition of claim 16 wherein the drug is selected from the group consisting of ceftriaxone, ketoconazole, ceftazidime, oxaprozin, albuterol sulfate, valacyclovir, urofollitropin, famciclovir, flutamide, enalapril, mefformin, itraconazole, buspirone, gabapentin, fosinopril, tramadol, acarbose, lorazepam, follitropin, glipizide, omeprazole, fluoxetine, lisinopril, levofloxacin, zafirlukast, interferon, growth hormone, interleukin, erythropoietin, granulocyte stimulating factor, nizatidine, bupropion, perindopril, erbumine, adenosine, alendronate, alprostadil, benazepril, betaxolol, bleomycin sulfate, dexfenfluramine, diltiazem, fentanyl, flecainid, gemcitabine, glatiramer acetate, granisetron, lamivudine, mangafodipir trisodium, mesalamine, metoprolol fumarate, metronidazole, miglitol, moexipril, monteleukast, octreotide acetate, olopatadine, paricalcitol, somatropin, sumatriptan succinate, tacrine, verapamil, nabumetone, trovafloxacin, dolasetron, zidovudine, finasteride, tobramycin, isradipine, tolcapone, enoxaparin, fluconazole, lansoprazole, terbinafine, pamidronate, didanosine, diclofenac, cisapride, venlafaxine, troglitazone, fluvastatin, losartan, imiglucerase, donepezil, olanzapine, valsartan, fexofenadine, calcitonin, and ipratropium.

18. The composition of claim 9 wherein the matrix further comprise an excipient selected from the group consisting of hydrophilic polymers, sugars, tonicity agents, pegylated excipients, and combinations thereof.

19. The composition of claim 9 wherein the mean diameter of the microparticles is between about 0.01 and 5  $\mu\text{m}$ .

20. The composition of claim 19 wherein the mean diameter of the microparticles is between about 1 and 5  $\mu\text{m}$ .

21. The composition of claim 9 wherein the microparticles are suspended in an aqueous solution suitable for parenteral administration.

22. The composition of claim 9 wherein the matrix is processed into

tablets or capsules suitable for oral administration.

23. The composition of claim 9 wherein the matrix is formed into suppositories suitable for vaginal or rectal administration.

24. The composition of claim 9 wherein the matrix is in a dry powder form suitable for pulmonary administration.

25. The composition of any of claims 9-24 wherein the porous matrix is made by a process comprising

(a) dissolving a drug in a volatile solvent to form a drug solution,

(b) combining at least one pore forming agent with the drug solution to form an emulsion, suspension, or second solution,

(c) incorporating at least one wetting agent into the emulsion, suspension, or second solution, and

(d) removing the volatile solvent and pore forming agent from the emulsion, suspension, or second solution to yield the porous matrix.

26. A method of delivering a drug to a patient in need thereof, comprising

administering a therapeutically or prophylactically effective amount of the drug in a formulation comprising the composition of any of claims 9-25.

27. The method of claim 26 wherein the formulation is suitable for administration by a route selected from the group consisting of parenteral, mucosal, oral, and topical administration.

28. The method of claim 27 wherein the parenteral route is selected from the group consisting of intravenous, intraarterial, intracardiac, intrathecal, intraosseous, intraarticular, intrasynovial, intracutaneous, subcutaneous, and intramuscular administration.

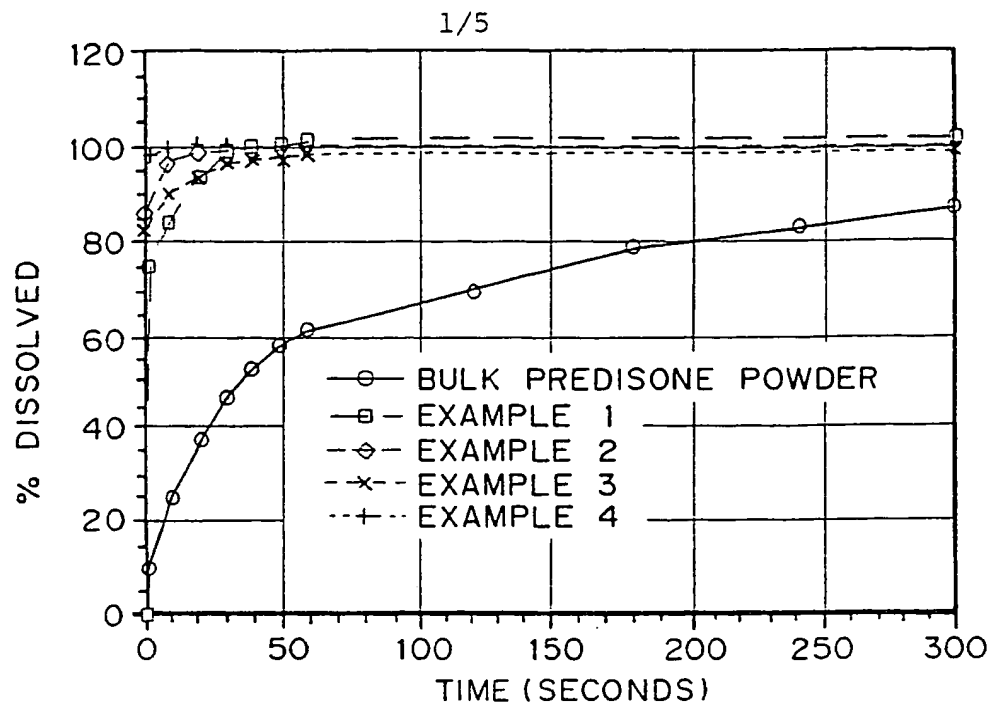
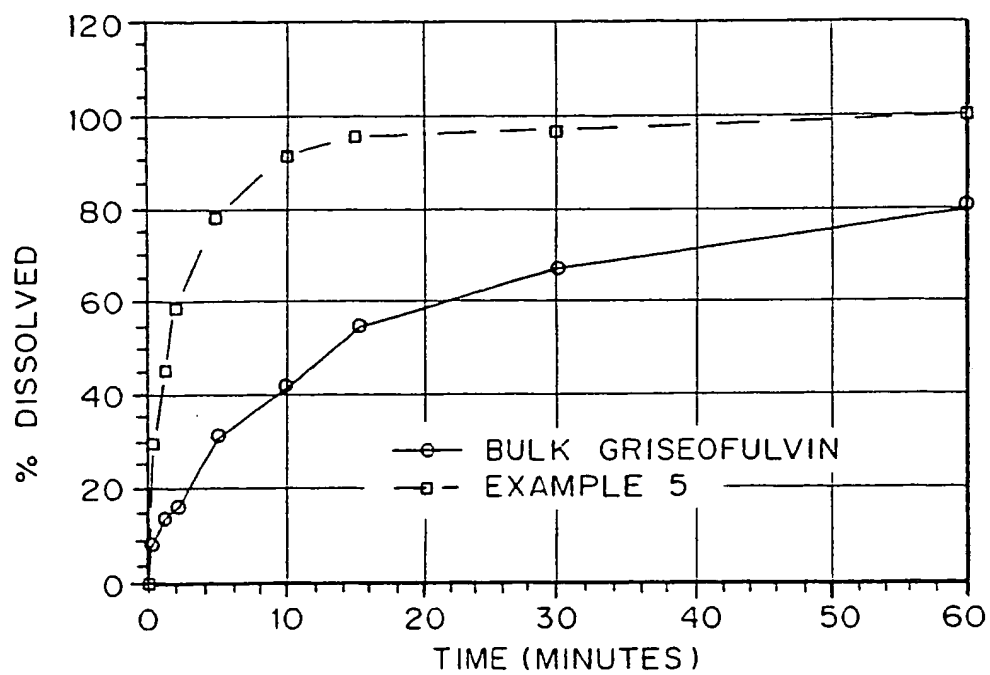
29. The method of claim 27 wherein the mucosal route is selected from the group consisting of pulmonary, buccal, sublingual, intranasal, rectal, and vaginal administration.

30. The method of claim 26 wherein the formulation is suitable for intraocular or conjunctival administration.

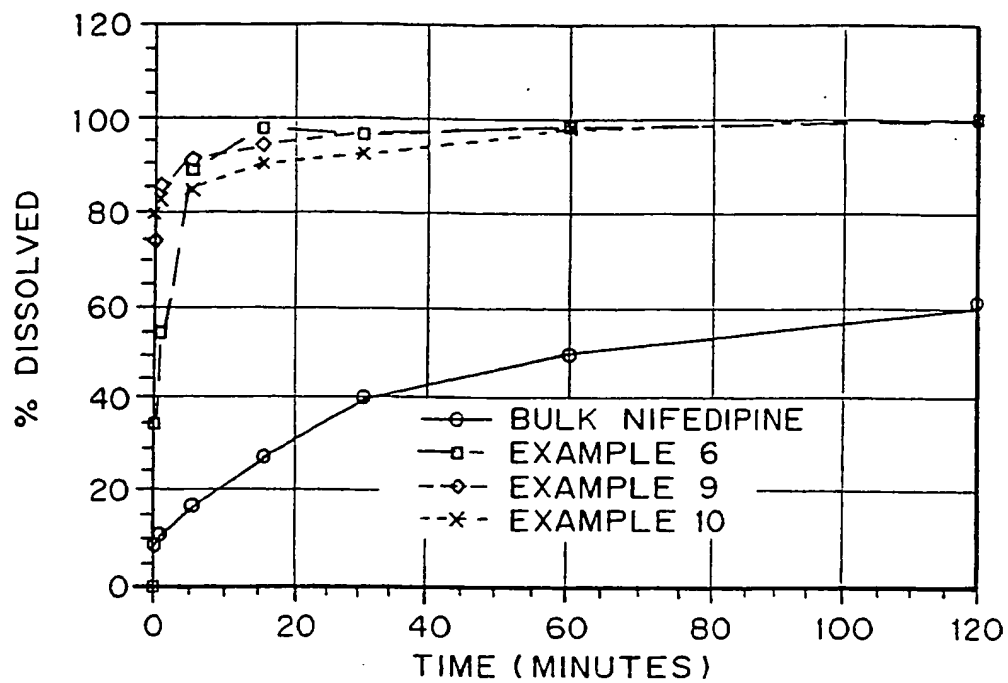
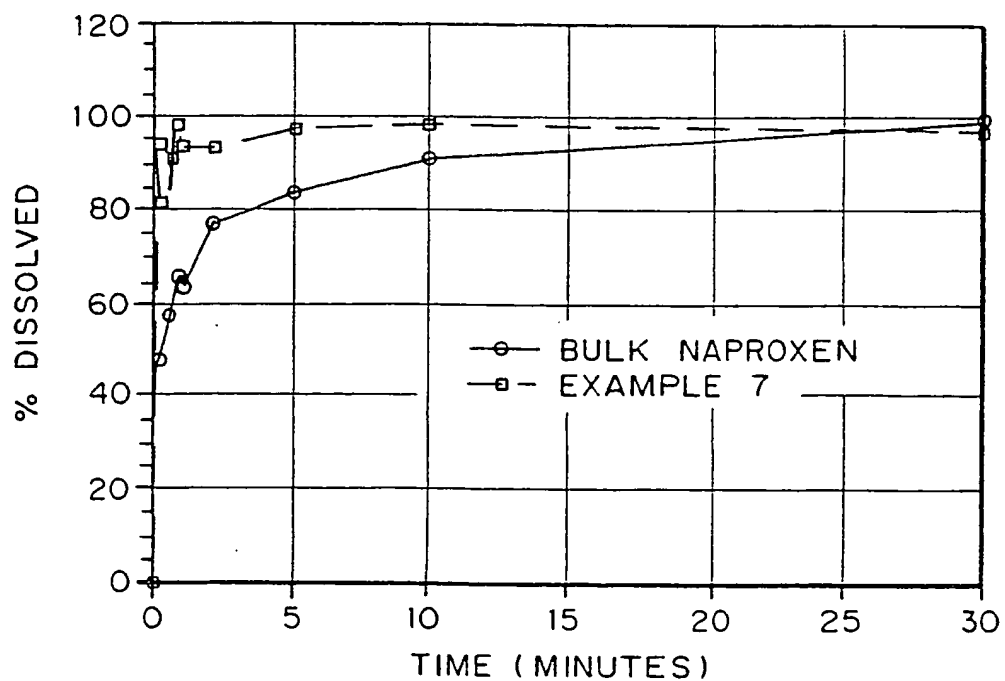
31. The method of claim 26 wherein the formulation is suitable for intracranial, intralesional, or intratumoral administration.

32. The method of claim 26 wherein the formulation is in an aqueous solution suitable for parenteral administration.

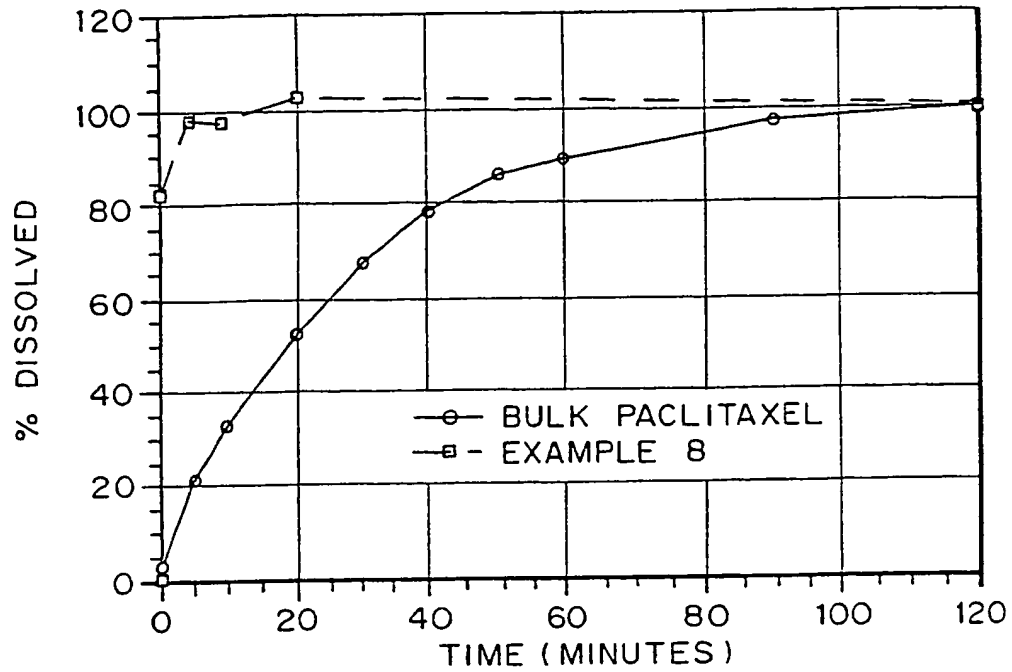
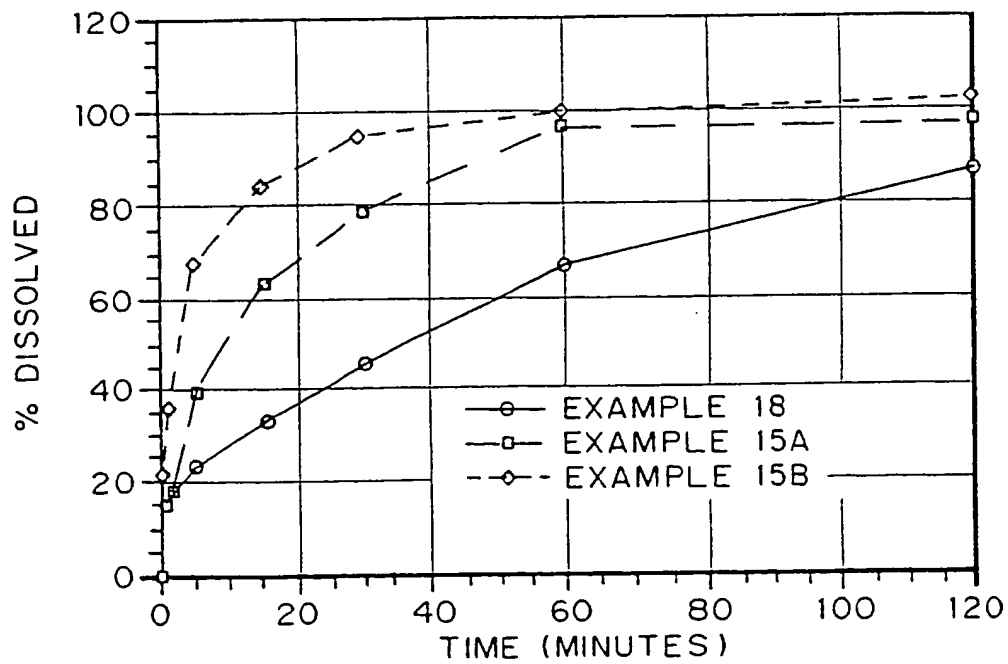


*FIG. 1**FIG. 2*

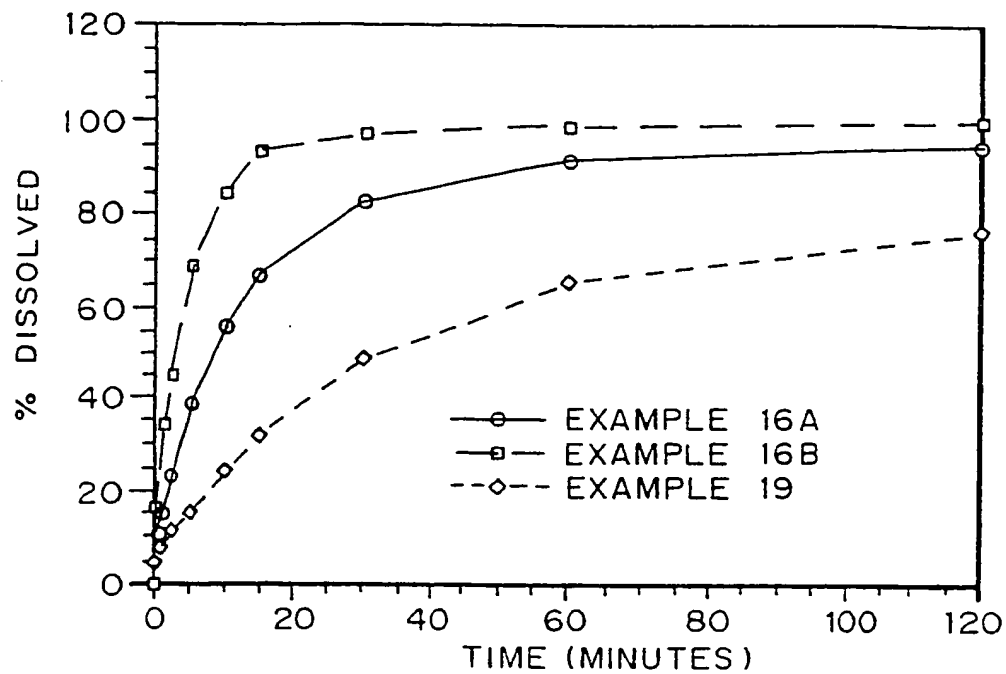
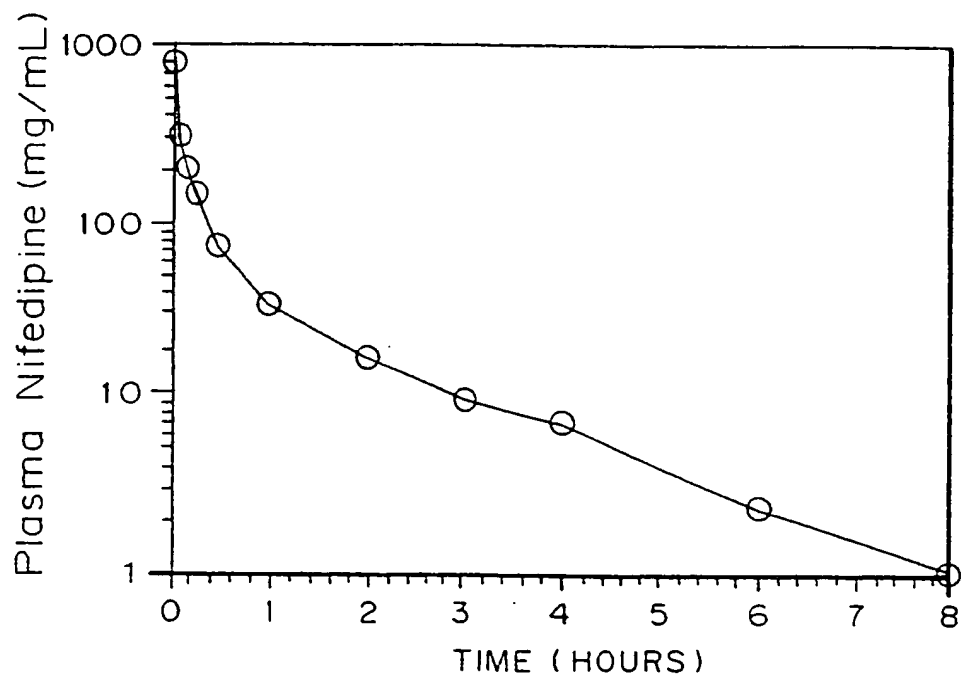
2/5

*FIG. 3**FIG. 4*

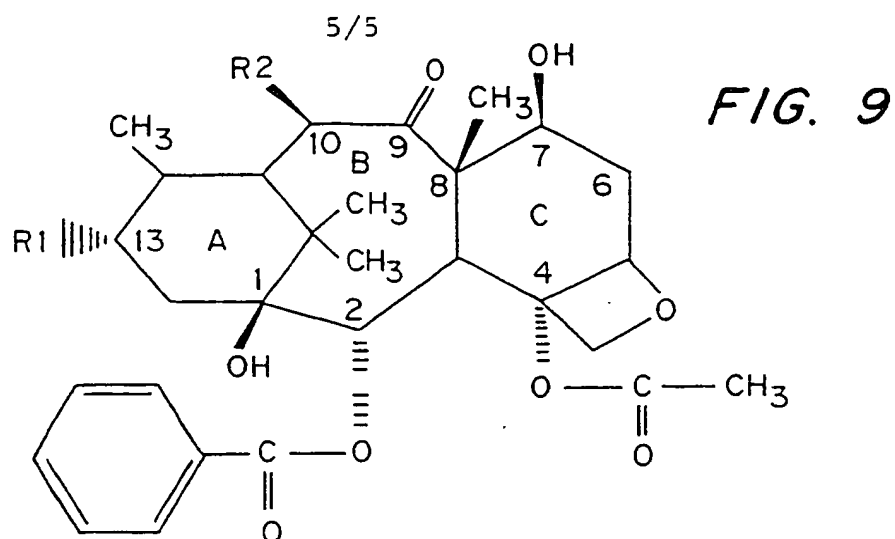
3/5

*FIG. 5**FIG. 6*

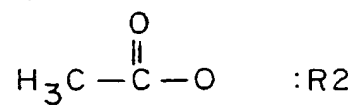
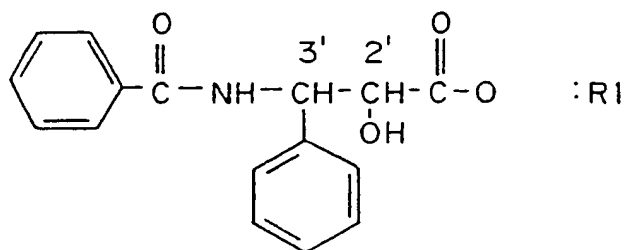
4/5

*FIG. 7**FIG. 8*

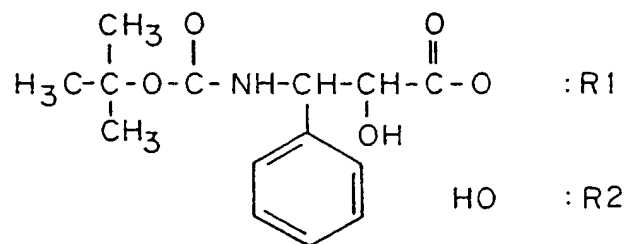
## Taxanes



PACLITAXEL :



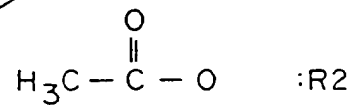
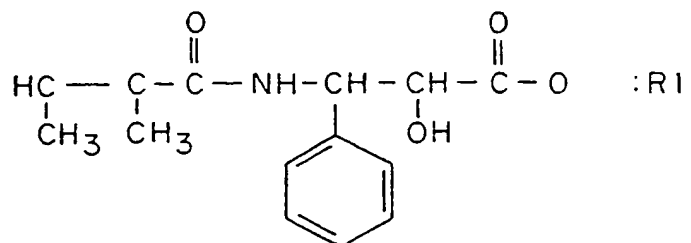
DOCETAXEL :



10-DEACETYLBACCATIN III:



CEPHALOMANNINE:



## INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/14578

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K9/16

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, PAJ, EPO-Internal

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 1 265 615 A (BOEHRINGER MANNHEIM G.M.B.H) 1 March 1972 (1972-03-01) the whole document ---	1, 3, 4, 6
X	DE 37 13 326 A (BASF AG) 29 October 1987 (1987-10-29) column 2, line 57 - column 5, line 15 ---	1-12, 18-20, 25
P, X, L	WO 99 56731 A (ACUSPHERE, INC.) 11 November 1999 (1999-11-11) the whole document page 3, line 15 - line 24 page 5, line 13 - line 14 page 14, line 16 - line 32 the priority claim of the present application might not be justified. --- -/--	1-32

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

## \* Special categories of cited documents:

\*A\* document defining the general state of the art which is not considered to be of particular relevance

\*E\* earlier document but published on or after the international filing date

\*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

\*O\* document referring to an oral disclosure, use, exhibition or other means

\*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*Z\* document member of the same patent family

Date of the actual completion of the international search

17 November 2000

Date of mailing of the international search report

24/11/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

Benz, K

# INTERNATIONAL SEARCH REPORT

Int .ional Application No

PCT/US 00/14573

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 655 237 A (HOECHST AKTIENGESELLSCHAFT) 31 May 1995 (1995-05-31)  the whole document ---	1-5, 8-12, 15, 18-20, 24-27, 29
X	WO 98 31346 A (MASSACHUSETTS INSTITUTE OF TECHNOLOGY ET AL.) 23 July 1998 (1998-07-23)  page 38; example 5 -----	1-5, 8-12, 15, 18-20, 24-27, 29

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/14578

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
GB 1265615	A	01-03-1972	NONE	
DE 3713326	A	29-10-1987	BE 1001516 A FR 2597719 A HU 44185 A, B IL 82245 A IT 1203957 B	21-11-1989 30-10-1987 29-02-1988 26-07-1990 23-02-1989
WO 9956731	A	11-11-1999	AU 2995499 A	23-11-1999
EP 655237	A	31-05-1995	AU 676390 B AU 7905194 A CA 2136704 A FI 945524 A HU 75152 A JP 7187996 A NO 944526 A NZ 264993 A ZA 9409378 A	06-03-1997 08-06-1995 28-05-1995 28-05-1995 28-04-1997 25-07-1995 29-05-1995 26-03-1996 11-08-1995
WO 9831346	A	23-07-1998	US 5855913 A EP 0954282 A US 5985309 A	05-01-1999 10-11-1999 16-11-1999